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## INSTRUCTIONS TO AUTHORS

1. MATERIAL should be original and not published elsewhere, in whole or in part.

2. LAYOUT should be as follows:

- (a) *Centred masthead to consist of*  
Title: informative but concise, without abbreviations and not including the names of new genera or species  
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Number of illustrations (figures, enumerated maps and tables, in this order)  
(b) *Abstract* of not more than 200 words, intelligible to the reader without reference to the text  
(c) *Table of contents* giving hierarchy of headings and subheadings  
(d) *Introduction*  
(e) *Subject-matter* of the paper, divided into sections to correspond with those given in table of contents  
(f) *Summary*, if paper is lengthy  
(g) *Acknowledgements*  
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3. MANUSCRIPT, to be submitted in triplicate, should be typewritten and neat, double spaced with 2.5 cm margins all round. First lines of paragraphs should be indented. Tables and a list of legends for illustrations should be typed separately, their positions indicated in the text. All pages should be numbered consecutively.

Major headings of the paper are centred capitals; first subheadings are shouldered small capitals; second subheadings are shouldered italics; third subheadings are indented, shouldered italics. Further subdivisions should be avoided, as also enumeration (never roman numerals) of headings and abbreviations.

Footnotes should be avoided unless they are short and essential.

Only generic and specific names should be underlined to indicate italics; all other marking up should be left to editor and publisher.

4. ILLUSTRATIONS should be reducible to a size not exceeding 12 × 18 cm (19 cm including legend); the reduction or enlargement required should be indicated; originals larger than 35 × 47 cm should not be submitted; photographs should be rectangular in shape and final size. A metric scale should appear with all illustrations, otherwise magnification or reduction should be given in the legend; if the latter, then the final reduction or enlargement should be taken into consideration.

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The number of the figure should be lightly marked in pencil on the back of each illustration.

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For books give title in italics, edition, volume number, place of publication, publisher.

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FISCHER, P.-H., DUVAL, M. & RAFFY, A. 1933. Études sur les échanges respiratoires des littorines. *Archs Zool. exp. gén.* 74: 627-634.  
KOHN, A. J. 1960a. Ecological notes on *Conus* (Mollusca: Gastropoda) in the Trincomalee region of Ceylon. *Ann. Mag. nat. Hist.* (13) 2: 309-320.  
KOHN, A. J. 1960b. Spawning behaviour, egg masses and larval development in *Conus* from the Indian Ocean. *Bull. Bingham oceanogr. Coll.* 17 (4): 1-51.  
THIELE, J. 1910. Mollusca: B. Polyplacophora, Gastropoda marina, Bivalvia. In: SCHULTZE, L. *Zoologische und anthropologische Ergebnisse einer Forschungsreise im westlichen und zentralen Süd-Afrika* 4: 269-270. Jena: Fischer. *Denschr. med.-naturw. Ges. Jena* 16: 269-270.

(continued inside back cover)

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LOWER CRETACEOUS  
(MIDDLE ALBIAN)  
AMMONITES FROM  
DOMBE GRANDE, ANGOLA

By  
MICHAEL R. COOPER

Cape Town      Kaapstad

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# LOWER CRETACEOUS (MIDDLE ALBIAN) AMMONITES FROM DOMBE GRANDE, ANGOLA

By

MICHAEL R. COOPER

*National Museum, Bulawayo*

(With 31 figures)

[MS accepted 3 March 1982]

## ABSTRACT

Two stratigraphically separated Middle Albian faunules, dominated by *Douvilleiceras* and *Oxytropidoceras* respectively, are described from the environs of Dombe Grande. The wide range of continuous variation within these assemblages has an important bearing on the taxonomy of these forms. The phylogeny of the Mojsisovicziinae is discussed, and the following new taxa are created: *Mortoniceratoides* gen. nov., *Oxytropidoceras* (*Mirapelia*) subgen. nov. and *O. (Benavidesites)* subgen. nov. Middle Albian biostratigraphy outside the Boreal Realm is discussed.

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## INTRODUCTION

Choffat (*in* Choffat & De Loriol 1888) was the first to record ammonites from the lower part of the Albian succession of Angola, describing and figuring an '*Acanthoceras*' *mamillare* (Schlotheim) (= *Douvilleiceras* *mammillatum* *aequinodum* (Quenstedt)) (Choffat & De Loriol 1888: 71, pl. 3 (fig. 1)) from the Dombe Grande area. The specimen was said to have come from the 'Couches à *Pholadomya pleuromyaeformis*', but as these beds are almost certainly of Aptian age (Howarth 1965; Cooper 1976) this is an error. Howarth (1965) described a small collection of *Douvilleiceras* made in 1930–1 by Alexandre Borges from the Dombe Grande region, and recorded the species *D. mammillatum* var. *?aequinodum* (Quenstedt) and *D. orbignyi* Hyatt, thereby suggesting the presence of upper Lower Albian (*D. mammillatum* Zone) strata. Cooper (1976) described the bio- and lithostratigraphy of the region. The present collections come from the immediate vicinity of Dombe Grande

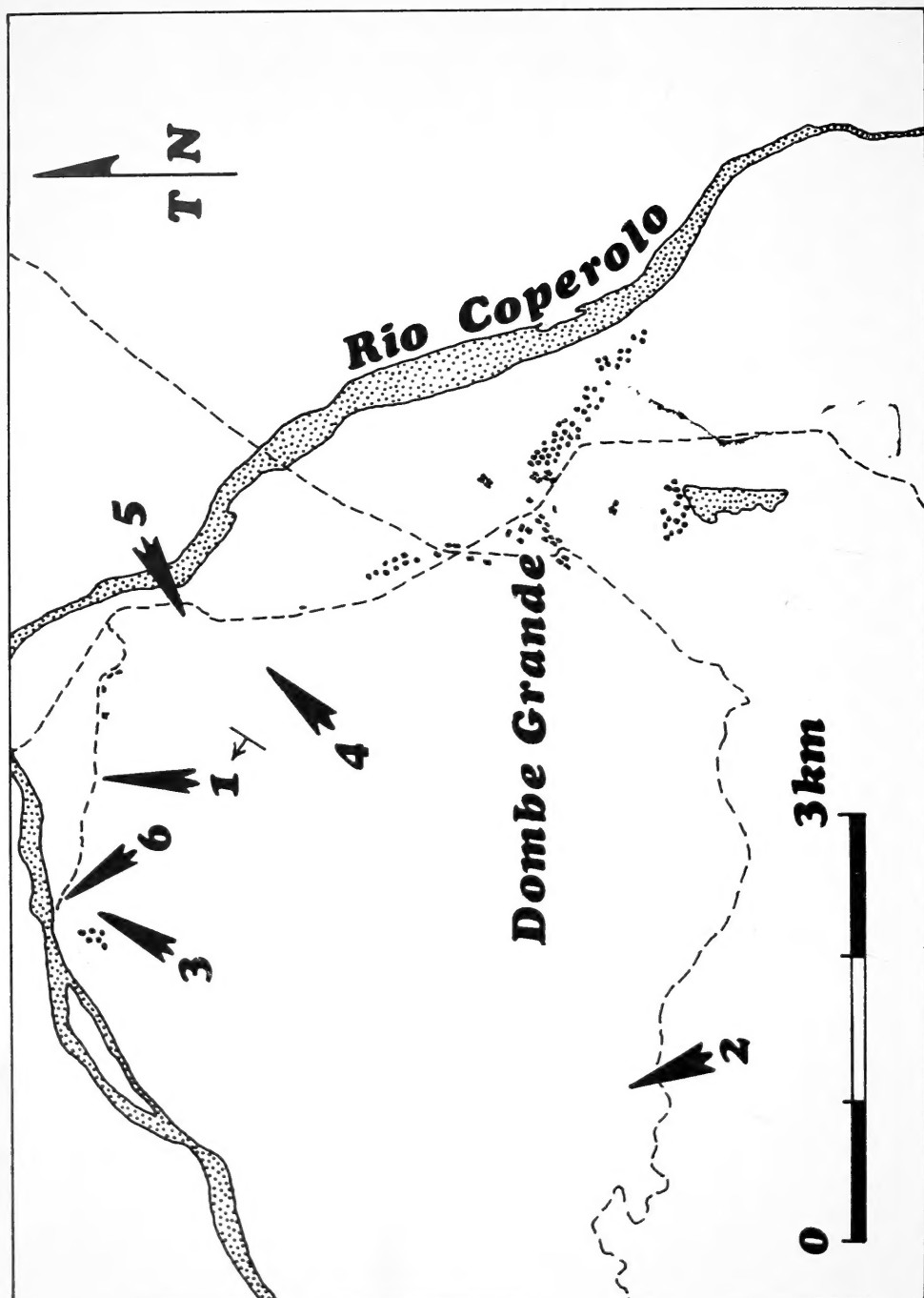


Fig. 1. Locality map, with collecting localities numbered. Broken lines represent roads.

(Fig. 1), and well above the beds containing *Pholadomya pleuromyaeformis* Choffat.

Prefixes to catalogue numbers refer to material in the following institutions:

BMNH	British Museum (Natural History), London
NHMP	Natural History Museum, Paris
OUM	Oxford University Museum, Oxford
SAM	South African Museum, Cape Town
USNMNH	U.S. National Museum of Natural History, Smithsonian Institution, Washington

Measurements are in millimetres, with dimensions, as a percentage of the diameter, in parentheses.

### REGIONAL GEOLOGY

The late Mesozoic succession in the Cuanza basin has been named the Cuanza Group (Cooper 1976). At Dombe Grande in the extreme south of the basin (12°54'S 13°13'E), the base of the Cuanza sequence is formed by the Dombe Formation, a unit of coarse clastics and lagoonal deposits with interbedded evaporites. The disconformably overlying transgressive deposits of the 'Formação com *Pholadomya*' (Neto 1960, 1961) attain a thickness of about 100 m and comprise a basal member of oolitic limestones, calcarenites and siltstones capped by calcareous siltstones rich in *Pholadomya pleuromyaeformis* and other bivalves (Fig. 2C–F). Also present at this level are the echinoids *Salenia dombeensis* De Loriol and *Pygurus africanus* De Loriol, and the molluscs *Neithea shawi* Pervinquière, *Retusa malheiroi* (Choffat), *Panopea gurgitis* (Brongniart), *Lithophaga praelonga* (d'Orbigny), *Nerita malheiroi* Choffat, '*Natica*' *feioi* Choffat, *Actaeonella* (*Trochactaeon*) *cordeiroi* (Choffat), *Actaeon lenzi* Choffat, *Ampullina* (*Pseudamaura*) *bulbiformis* (J. Sowerby), *Pinna robinaldina* d'Orbigny, *Cylindrites?* *delgadoi* Choffat, and *Tylostoma peschueli* Choffat.

Marine regression is reflected in the succeeding 'Formação com *Nerinea*' (Fig. 1, locality 2), a lithologically highly variable unit spanning some 400 m and comprising red and yellow lagoonal clastics yielding fossil wood, the gastropods *Nerinea capelloi* Choffat (Fig. 3G–H) and *Cerithium monteiroi* Choffat, and an indeterminate ammonite (*vide* Neto 1961).

The 'Formação com *Nerinea*' are overlain, perhaps disconformably, by limestones and marls at the base of the Cuio Formation, the lower beds of which have yielded the faunas described herein. The two assemblages, which were picked up mainly as surface scree, are stratigraphically separated at locality 4 (Fig. 1), with the *Oxytropidoceras* fauna occurring to the north-east and thus from the younger beds. Upward the sequence becomes increasingly unfossiliferous, and calcarenites and grits (Fig. 1, locality 1) span much of the

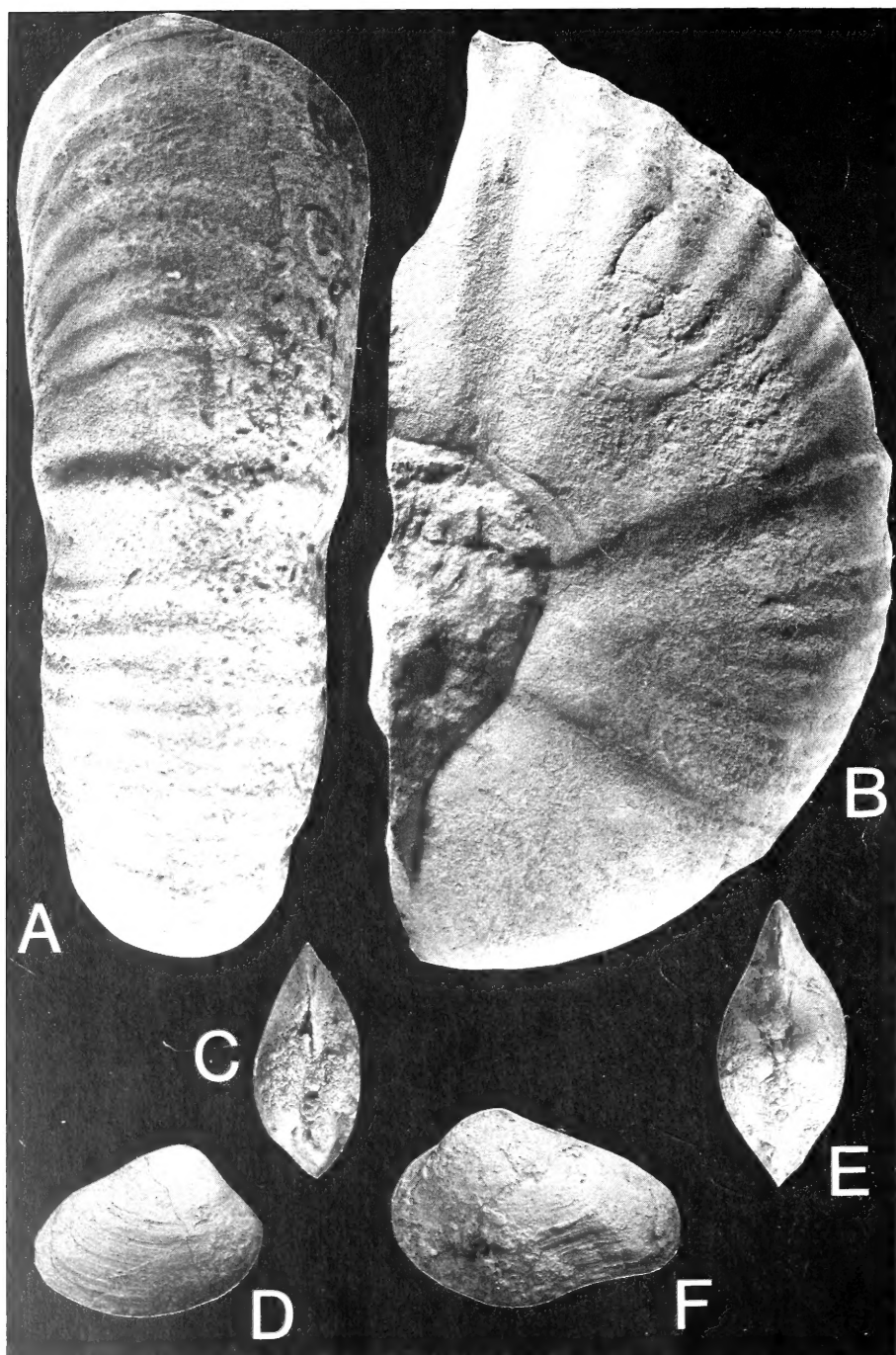


Fig. 2. A-B. *Puzosia bistricta* (White). SAM-PCA2649.  $\times 1$ . C-F. Undetermined bivalves from the 'Couches a *Pholadomya*', Reinecke Collection, South African Museum,  $\times 1.5$ .



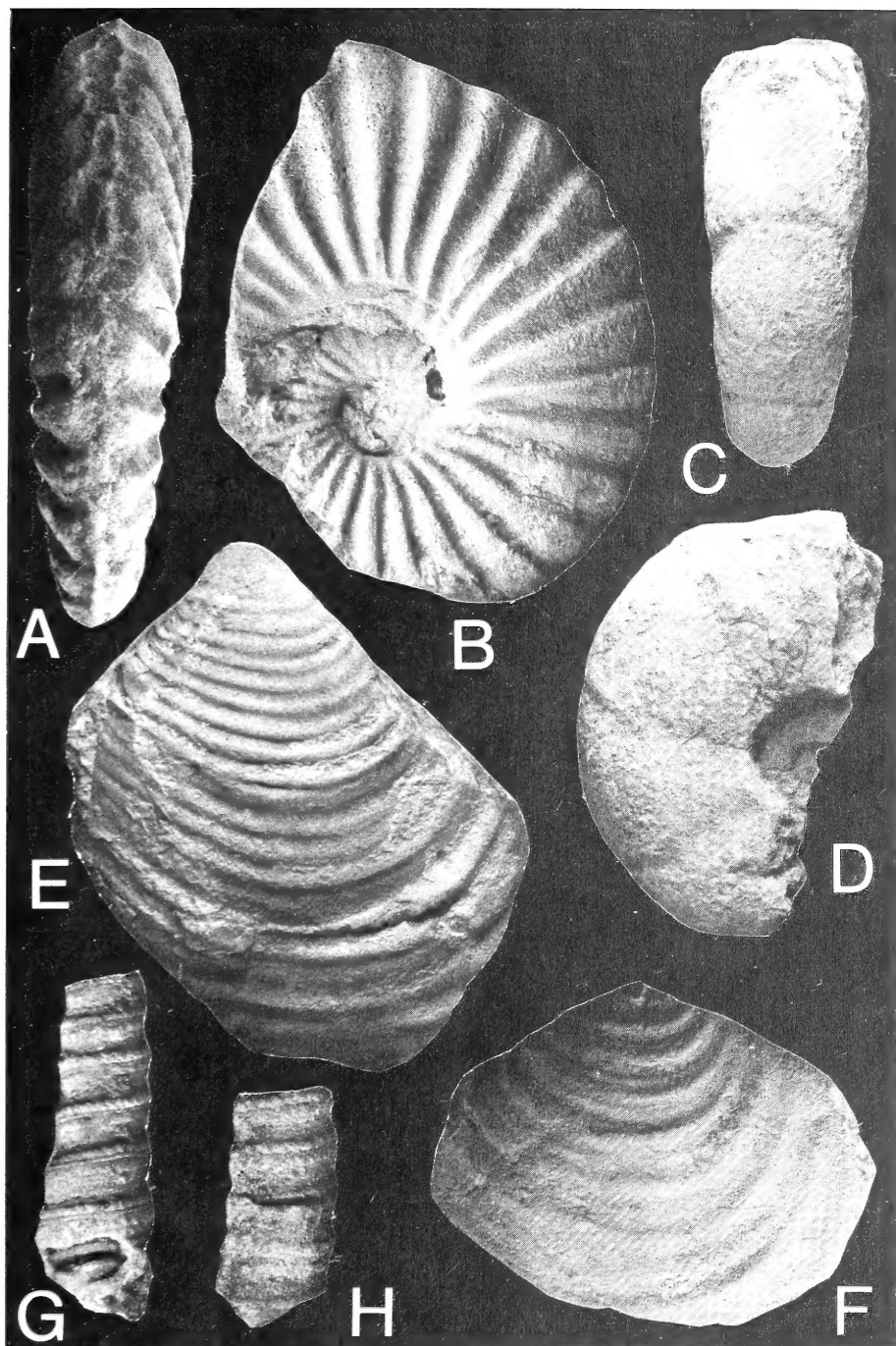


Fig. 3. A-B. *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White), SAM-PCA5445, an individual transitional to *douglasi* morphotypes. C-D. *Puzosia bistricta* (White), SAM-PCA3474. E. *Inoceramus* gr. *anglicus* Woods, SAM-PCA3443, from the *Douvilleiceras* assemblage. F. *Birostrina?* cf. *coptensis* (Casey), SAM-PCA2664, from the *Oxytropidoceras* assemblage. G-H. *Nerinea capelloi* Hoffat. G. SAM-PCA5448,  $\times 1.5$ . H. SAM-PCA5450,  $\times 1.5$ . All  $\times 1$  unless otherwise stated.

remainder of the Middle Albian. These sediments are interpreted as reflecting marine regression. The Cuio Formation attains a thickness of some 200 m and is succeeded by bioclastic and oolitic limestones and calcarenites (Fig. 1, localities 3, 6) yielding a low Upper Albian fauna including *Prohysterocheras wordiei* Spath and *Mortoniceras* spp. This unit is to be correlated with the Catumbela Member of the Cabo Ledo Formation to the north, reflecting the early late Albian (*Hysterocheras orbignyi* Zone) transgression.

## SYSTEMATICS

Subclass AMMONOIDEA Zittel, 1884

Family **Douvilleiceratidae** Parona & Bonarelli, 1897

### *Discussion*

Since the lineage *Procheloniceras* → *Cheloniceras* → *Eodouvilleiceras* → *Douvilleiceras* represents a single phyletic lineage, the subfamily Cheloniceratinae should be included in the synonymy of the nominate subfamily.

Subfamily **Douvilleiceratinae** Parona & Bonarelli, 1897

Genus *Douvilleiceras* de Grossouvre, 1894

Type species *Ammonites mammillatus* Schlotheim, 1813; subsequent designation of I.C.Z.N., Opinion 422, 1956

### *Discussion*

Casey (1962: 260) has given a detailed diagnosis and discussion of this genus, to which little may be added.

The genus *Trinitoceras* was erected (Scott 1940: 1016) for very large douvilleiceratids in which the '... douvilleiceratid sculpture is evident up to or beyond a diameter of 200 mm. Costae are multituberculate as in *Douvilleiceras*, but the umbilical tubercles are greatly exaggerated in size, forming large bullae on the umbilical margin at the ends of alternate costae. On later whorls costae lose all trace of tuberculation but retain, irregularly, their alternate primary and secondary arrangement. On the latest whorls shown, the ribs are nearly equal, except that the occasional secondary (marked only by the fact that it does not reach the umbilicus) is intercalated between the primaries. The whorl section, degree of evolution, and the umbilical area do not differ greatly from like features of well-known species of *Douvilleiceras*.'

It is clear from the above description that the genus *Trinitoceras* is based upon the gigantic shells that accompany virtually all *Douvilleiceras* populations; thus *D. leightonense* Casey, *D. cheloniceratiforme* Tavani, *D. restitutum* Anderson, and *D. charshangense* Mirzoyev are all typical *Trinitoceras*. However, the fact that *Trinitoceras* and *Douvilleiceras* everywhere occur together as well as the fact that they are indistinguishable in their early ontogenetic stages and

attain maturity at vastly different diameters, suggests to the writer that the differences are those between sexual dimorphs. *Trinitoceras* is here considered to be based upon *Douvilleiceras* macroconchs, and Casey (1962) is followed in regarding it a synonym of *Douvilleiceras*.

Up to now almost fifty trivial names have been applied to *Douvilleiceras*; these include *D. mammillatum* (Schlotheim) and its varieties *paucicostatum* Parona & Bonarelli, *praecox* Casey, *baylei* Spath and *aequinodum* (Quenstedt), *D. inaequinodum* (Quenstedt), *D. pseudinaequinodum* Collignon, *D. orbignyi* Hyatt, *D. clementinum* (d'Orbigny), *D. solitae* (d'Orbigny), *D. tarapacaense* Etayo-Serna, *D. abozaglio* Etayo-Serna, *D. ivernoisi* (Coquand), *D. spiniferum* (Whiteaves), *D. rex* (Scott), *D. reesidei* (Scott), *D. grandense* (Scott), *D. adkinsi* (Scott), *D. dunlapi* (Scott), *D. quitmanense* Scott, *D. spathi* Scott, *D. cuchillense* Scott, *D. muralense* Stoyanow, *D. aurarium* Anderson, *D. restitutum* Anderson, *D. offarcinatum* (White), *D. euzebioi* (Maury), *D. sergipense* Beurlen, *D. multinodosum* Hubach, *D. benonae* Besairie, *D. spinosum* Tavani, *D. albiense* (Tavani), *D. variabile* Tavani, *D. cheloniceratiforme* Tavani, *D. scabrosum* Casey, *D. alternans* Casey, *D. pustulosum* Casey, *D. leightonense* Casey and its variety *pringlei* Casey, *D. magnodosum* Casey, *D. subleightonense* Mirzoyev, and *D. charshangense* Mirzoyev.

The extreme range of morphological variation exhibited by most assemblages is emphasized by the splitting of the British collections into fourteen taxa (Casey 1962), those from Texas into eleven species (Scott 1940), and the Somali collections into eight species (Tavani 1942, 1949), and this seems to suggest genetic polymorphism. Thus Casey (1962: 263) has noted that all the mostly endemic species of *Douvilleiceras* recorded from southern England '... come from a limited range of strata and they constitute a closely interconnected plexus that in former days would have been interpreted (perhaps correctly) as one very large, highly variable species'. Unfortunately Casey's revision, coming at a time when it was the practice to base species on very narrowly defined limits, suffers from oversplitting, since recent studies of ammonite populations (Kennedy & Cobban 1976) suggests that such variation is not extraordinary. In addition, Mirzoyev (1967: 54) has also commented that '... all species of *Douvilleiceras* are closely interrelated and could have been treated as a single very extensive and varied species; but investigators concerned with this group of ammonites have made efforts to discover differences, based entirely on the frequency, degree of development and modifications of the conch sculpture in the course of its development'.

Casey (1962), in his revision of the British species of *Douvilleiceras*, laid much emphasis on the width of the ventral sulcus, number of notches per rib, and the ontogenetic stage at which ornament change occurred as taxonomic criteria. However, many writers have noted that the number of notches increases with diameter, while Collignon (1963), Mirzoyev (1967), and McLearn (1972) have all reported specimens with different numbers of notches on opposing flanks. Consequently, this criterion is considered to be of very

limited taxonomic significance, if at all. Both the width of the ventral sulcus and the ontogenetic stage at which morphological changes occur are relative characters that are unlikely to stand up to population analysis, especially when it is considered that many of the taxa so defined are contemporaneous, occurring together in the same bed. As such, the writer believes *Douvilleiceras* to display genetic polymorphism and thus high intraspecific variability.

Stratigraphically *Douvilleiceras* ranges from the top of the *Leymeriella tardefurcata* Zone (*Leymeriella regularis* Subzone) into the lower part of the Middle Albian (*Lyelliceras lyelli* Subzone of the *Hoplites dentatus* Zone). During this period it attained a cosmopolitan distribution and is currently reported from North America (British Columbia, California, Arizona, Texas, New Mexico), South America (Peru, Columbia, Brazil), Africa (Gabon, Angola, Zululand, Mozambique, Somalia), Madagascar, Asia (eastern Carpathians, northern Caucasus, western and central Kopet Dag, Bol'shoy Balkhan, Tuarkyr, Mangyshlak, Badkhiz Islands, southern Gissar Range, Afghanistan), Europe (southern England, France, Switzerland, ?Germany, Poland, Bulgaria), and Japan.

*Douvilleiceras mammillatum aequinodum* (Quenstedt, 1849)

Figs 4A–B, 6C–G, 7D, G–I, 8A–B, 9A–C, E, 10C–D, F–H, 12A–B, 17A

*Ammonites monile aequinodus* Quenstedt, 1849: 137, pl. 10 (fig. 2).

? *Ammonites offarcinatus* White, 1887: 219, pl. 23 (fig. 4 only).

*Acanthoceras mammillare* (Schlotheim) Choffat (in Choffat & De Loriol), 1888: 71, pl. 3 (fig. 1).

*Douvilleiceras monile* var. *aequinodum* (Quenstedt) Breistroffer, 1947: 65.

? *Douvilleiceras cheloniceratiforme* Tavani, 1949: 38, pl. 8 (fig. 6).

? *Douvilleiceras monile* (J. Sowerby) Benavides-Cáceres, 1956: 442, pl. 43 (figs 1–4).

? *Douvilleiceras restitutum* Anderson, 1938: 175, pl. 54 (fig. 2).

*Douvilleiceras mammillatum* var. *aequinodum* (Quenstedt) Casey, 1962: 271, pl. 40 (fig. 5), pl. 41 (figs 5–7), pl. 42 (fig. 10), figs 94a–c, 95a–b, 102d, 103a–b. Howarth, 1965: 343, pl. 1 (figs 1–4).

? *Douvilleiceras albense* Spath, Collignon, 1963: 112, pl. 284 (fig. 1243).

*Douvilleiceras* sp. B, Beurlen, 1970: 460, pl. 4 (figs 3–4), figs 16–17.

? *Douvilleiceras spiniferum* (Whiteaves) McLearn, 1972: 62, pl. 28 (fig. 1 only).

*Douvilleiceras* cf. *mammillatum* (Schlotheim) Young, 1974: 184, pl. 4 (figs 1, 3, 6).

### Material

Fifty-seven specimens, SAM-PCA2639, 3409, 3417, 3427, 3429–3431, 3434, 3439–3440, 3442, 3456, 3459–3460, 3462–3464, 3467, 3469, 3471–3473, 3476–3480, 3482, 3484–3487, 3490–3491, 3493–3494, 3499, 3501, 3503, 3509–3511, 3516–3518, 3520, 3522–3523, 3526–3527, 3529–3530, 3532, 3534, and 3536, all preserved as limestone steinkerns.

### Description

This taxon is the most abundant component of the *Douvilleiceras* fauna, although many of the specimens are fragmentary and most are more or less eroded, having been picked up as surface scree. This erosion gives a number of

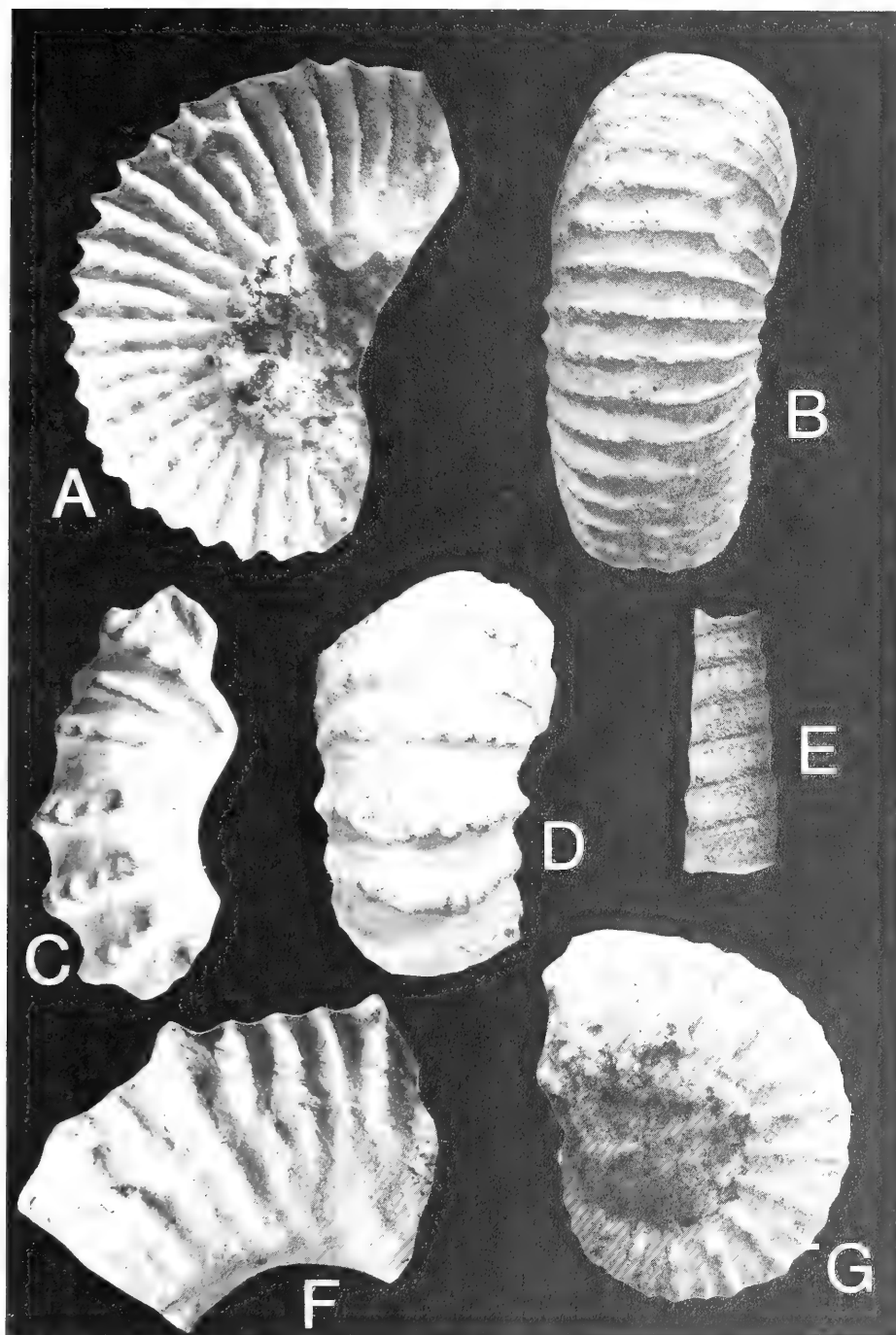


Fig. 4. A-B. *Douvilleiceras mammillatum aequinodum* (Quenstedt), SAM-PCA3434. C-D, F. *Douvilleiceras inaequinodum* (Quenstedt). C-D. SAM-PCA3435, a juvenile showing very fine intercalated ribs. F. SAM-PCA5451, a badly eroded fragment showing three intercalated ribs between main ribs. E. *Nerinea capelloi* Choffat, SAM-PCA5449,  $\times 1.5$ . G. *Douvilleiceras variabile* Tavani, SAM-PCA3448. All  $\times 1$  unless otherwise stated.

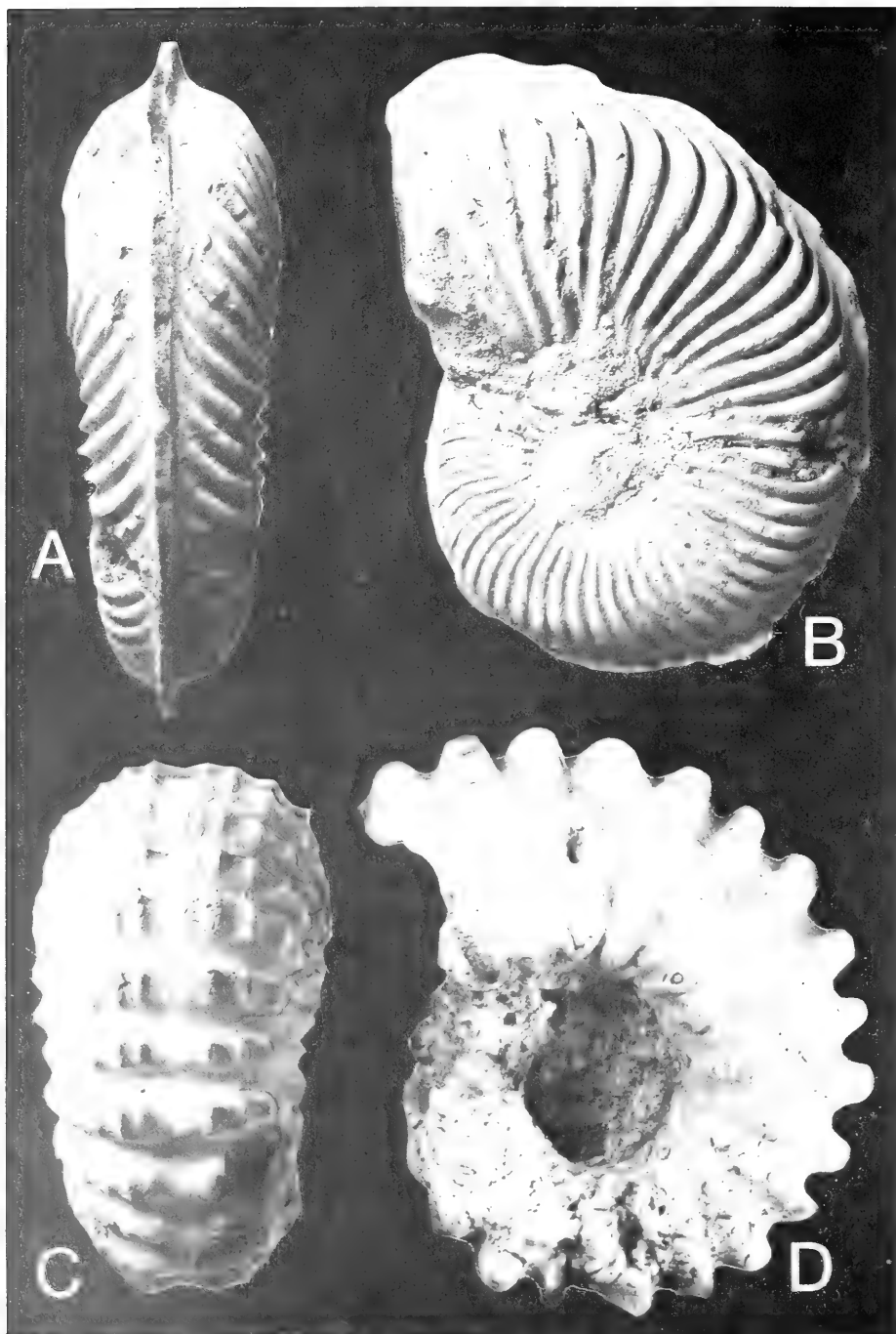


Fig. 5. A-B. *Oxytropidoceras* (*Oxytropidoceras*) *roissyanum* (d'Orbigny). The original of D'Orbigny's (1841) photograph, NHMP. C-D. *Douvilleiceras mammillatum mammillatum* (Schlotheim). The neotype, BM-C12491. Both  $\times 1$ . Photos: W. J. Kennedy.

specimens the appearance of having had very much narrower ribs than, in fact, they probably had.

The shell is moderately evolute (umbilicus 32,5–39% of the diameter), with rather inflated whorls, and a subreniform depressed whorl section. In all the better preserved material the ventral sulcus is, or appears to have been, rather narrow. The number of notches to each rib varies with ontogeny, with about 5 at 30 mm diameter increasing to 8–9 at about 60 mm diameter. Up to about 40 mm diameter the ribs tend to be simple, distant and rursiradate, commonly with 10–14 ribs per half whorl. However, several of the more densicostate juveniles have 16 ribs per half whorl. Beyond this diameter intercalation and bifurcation becomes increasingly common, generally with 16–20 rather uniform ribs per half whorl. One individual, SAM-PCA3417, retains simple ribbing to 62 mm diameter, whereas SAM-PCA3431 has very fine, nontuberculate intercalatories between main ribs at about 40 mm diameter, only to lose them with further growth. The tuberculation of this subspecies is very reduced, typically with the lateral tubercle subdued and bullate. In SAM-PCA3472, however, main ribs bear a prominent lateral tubercle and, up to 35 mm diameter, alternate with fine nontuberculate intercalatories. Beyond this diameter all the ribbing becomes uniform and the lateral tubercle is much reduced and bullate. In SAM-PCA3435, the early whorls are strongly tuberculate and distantly ribbed but, at 47 mm diameter, there is a sudden and dramatic reversion to normal *aequinodum*-type ornament. SAM-PCA3430 connects typical *aequinodum*-type morphologies to the large macroconch fragments (SAM-PCA3408, 3410–3411, 3414, 3425, 3433, 3465, 3483, 3500) in the collection.

#### Measurements

No	D	H	W	W/H	U
SAM-PCA3427	75	29	34	1,17	29 (39)
SAM-PCA3429	83	35	45	1,29	27 (32,5)
SAM-PCA3431	?	32	37	1,16	?
SAM-PCA3434	70	26	32	1,23	27 (38,5)
SAM-PCA3527	51	19	21	1,11	18 (35,3)

#### Discussion

Quenstedt (1849) described two species of *Douvilleiceras*, *D. aequinodum* and *D. inaequinodum*. Since the latter species is typically lower Middle Albian (Casey 1962), it may be reasonable to assume this is also the level of *D. aequinodum*. Such a supposition tends to be supported by their coexistence in the *eodontatus* Nodule Bed (Bed e of Destombes *et al.* 1977) (Fig. 12A–D, G–H) at Bully, Pays de Bray, France, and also in Angola. Since the Angolan *D. mammillatum* material is dominated by *aequinodum* morphotypes, with no undoubted individuals of *D. mammillatum* s.s. (Fig. 5C–D), it seems reasonable to assume that a genuine shift in population structure has occurred when



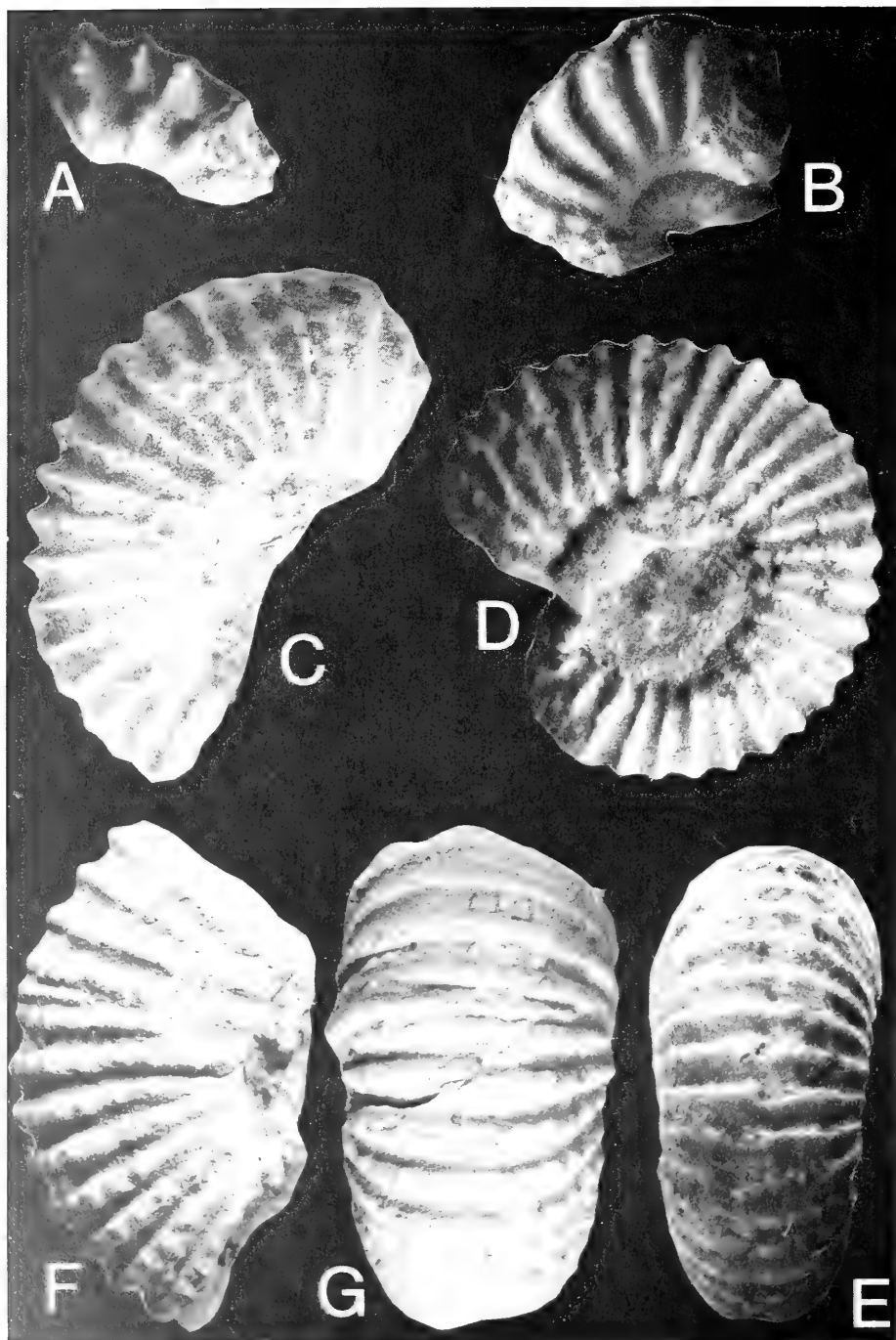


Fig. 6. A. *Douvilleiceras inaequinodum* (Quenstedt), SAM-PCA3535, a juvenile. B. *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White), SAM-PCA3765, a typical juvenile. C-G. *Douvilleiceras mammillatum aequinodum* (Quenstedt). C. SAM-PCA3427. D-E. SAM-PCA2639. F-G. SAM-PCA3431. All  $\times 1$ .



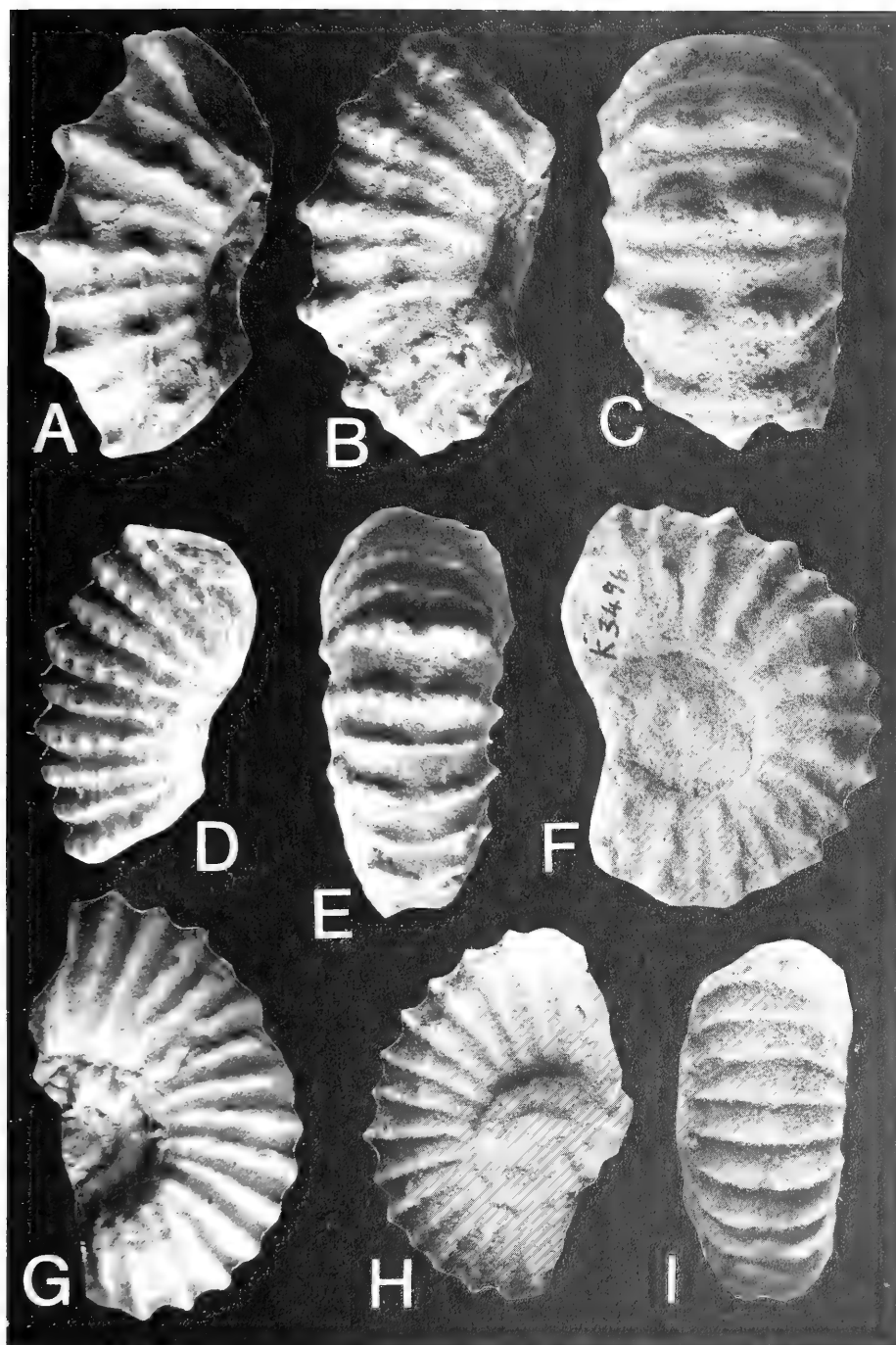


Fig. 7. A-C. *Douvilleiceras inaequinodum* (Quenstedt). A. SAM-PCA3466. B-C. SAM-PCA3437, a typical example. D, G-I. *Douvilleiceras mammillatum aequinodum* (Quenstedt). D. SAM-PCA3479. G. SAM-PCA3481. H-I. SAM-PCA3527, a typical juvenile with fine, distant, simple ribs. E-F. *Douvilleiceras variabile* Tavani, SAM-PCA3496. All  $\times 1$ .

compared with the typical *mammillatum* Zone assemblages of *D. mammillatum* (cf. Casey 1962). It is for this reason, and because *D. aequinodum* can arguably be regarded as typically Middle Albian, that the writer has used Quenstedt's name at subspecific level. As noted by Casey (1962) *aequinodum* morphotypes are also abundant in the *mammillatum* Zone of England, but at this level coexist with, and are connected by transitions to, *D. mammillatum* s.s., *D. mammillatum* var. *praecox* Casey, and *D. monile* (J. Sowerby). Since there is, in the writer's opinion, but a single biological taxon involved in the latter list, the various nominal forms are merely morphotypes within a single polytypic species. Disregarding the nomenclatural problems, it is clear that the Angolan population of *D. mammillatum* is substantially different from that at Copt Point, Folkestone, and thus warrants subspecific distinction.

Both *Douvilleiceras restitutum* Anderson and *D. cheloniceratiforme* are based upon macroconchs whose inner whorls suggest they may belong here. Similarly, the hypotype of *D. spiniferum* (Whiteaves) figured by McLearn (1972, pl. 28 (fig. 1)) also shows few features to distinguish it from typical *aequinodum* morphotypes. In addition, one of White's (1887, pl. 23 (fig. 4)) syntypes of *D. offarcinatum* shows a rib density that suggests possible reference to Quenstedt's taxon.

#### *Douvilleiceras variabile* Tavani, 1949

Figs 4G, 7E-F, 8C-D, 9G, 11C-D, 13A, 15E, 22A

*Ammonites mammillaris* Schlotheim, d'Orbigny, 1841: 249, pl. 72 (fig. 5 only).

*Douvilleiceras variabile* Tavani, 1949: 37, pl. 10 (figs 1-2).

? *Douvilleiceras pseudinaequinodum* Collignon, 1963: 120, pl. 288 (fig. 1249).

*Douvilleiceras monile* (J. Sowerby) Collignon, 1963: 118, pl. 287 (fig. 1248).

*Douvilleiceras* aff. *baylei* Spath, Collignon, 1963: 118, pl. 287 (fig. 1247).

*Douvilleiceras subleightonense* Mirzoyev, 1967: 59, pl. 8 (figs 1-3).

? *Douvilleiceras* sp. B, McLearn, 1972: 67, pl. 13 (fig. 1), pl. 16 (fig. 1).

#### Material

Four specimens, SAM-PCA3422, 3447-3448 and 3496, all preserved as internal moulds.

#### Description

In shell form, this species is very similar to the last but with substantially different ornament. In the Angolan material, ornament comprises well-developed main ribs ornamented with a distinct umbilical tubercle, a pronounced mid-lateral tubercle and bullate ventral tubercles that weaken beyond about 45 mm diameter and are lost at about 60 mm diameter. At about this stage a small tubercle also appears between the umbilical and mid-lateral tubercle, and the ventral tubercles bear three notches. There are, therefore, eight notches per rib. Between these long ribs are invariably one, sometimes two intercalatories that do not connect with the umbilicus and lack mid-lateral tubercles. These ribs tend to be somewhat finer than the main ribs and give rise to the typical 'variable' ornament of this species.

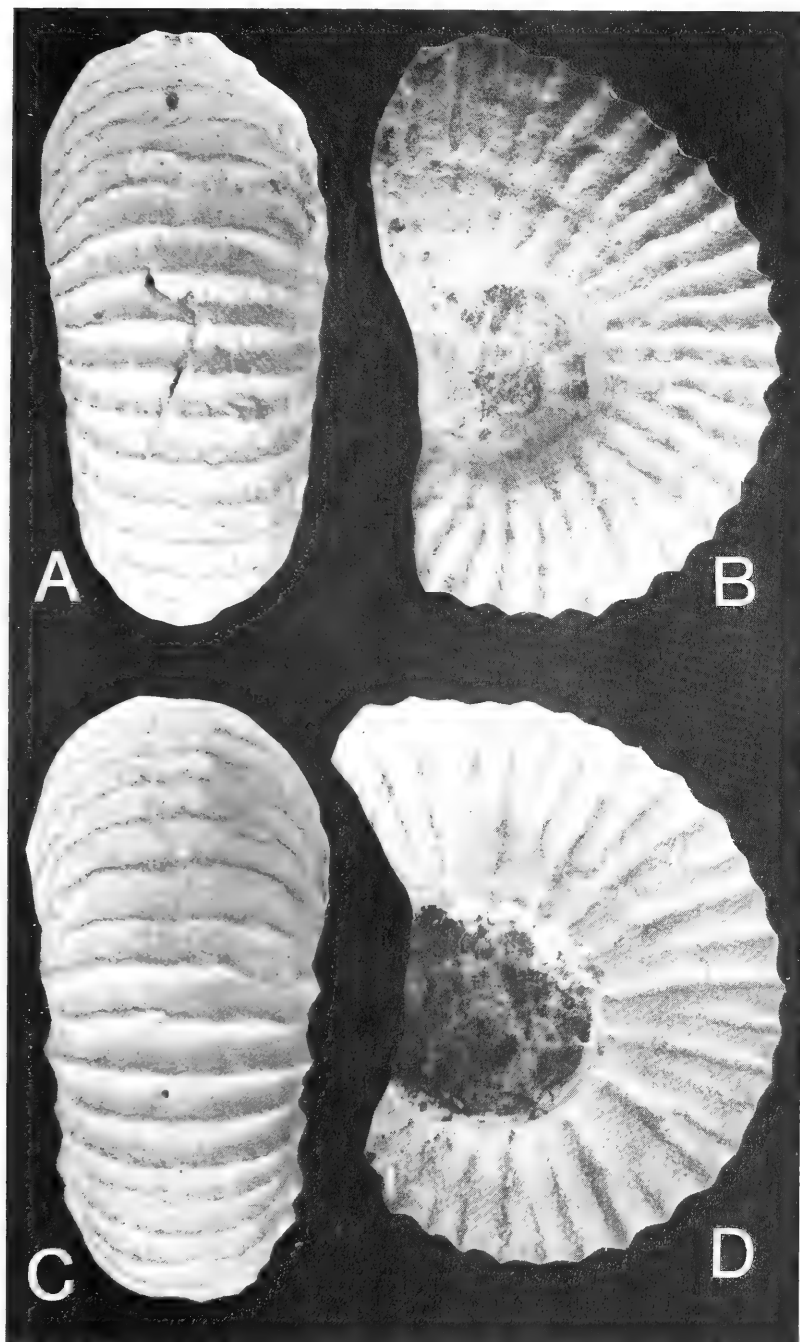


Fig. 8. A-B. *Douvilleiceras mammillatum aequinodum* (Quenstedt), SAM-PCA 3429. C-D. *Douvilleiceras variabile* Tavani, SAM-PCA3447. Note the alternating long and short ribs and prominent lateral tubercle typical of this species. Both  $\times 1$ .

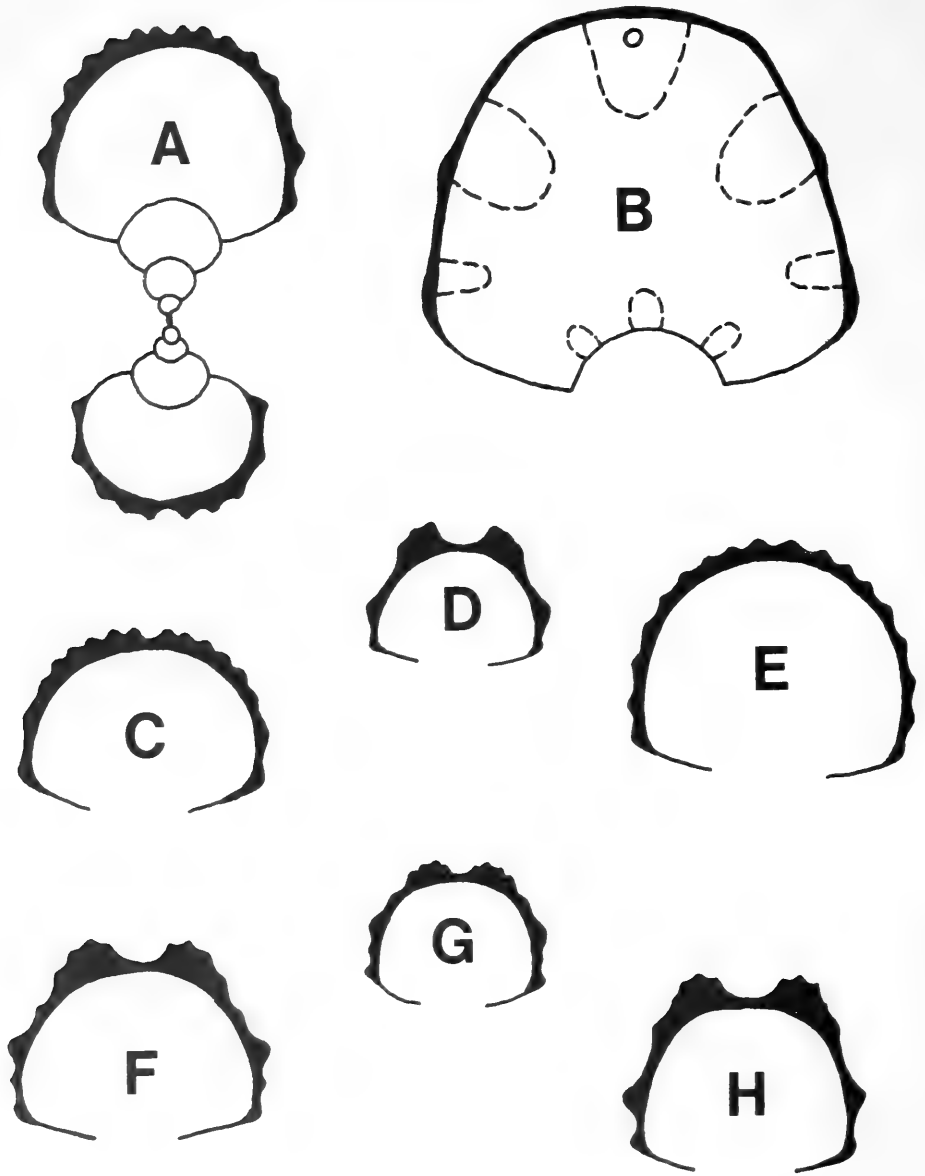


Fig. 9. Whorl sections of *Douvilleiceras*. A-C, E. *Douvilleiceras mammillatum aequinodum* (Quenstedt). A. SAM-PCA3417. B. SAM-PCA3433. C. SAM-PCA3479. E. SAM-PCA 3431. D, F, H. *Douvilleiceras inaequinodum* (Quenstedt). D. SAM-PCA3535. F. SAM-PCA3437. H. SAM-PCA 3466. G. *Douvilleiceras variabile* Tavani. SAM-PCA3496. All  $\times 1$ .

*Measurements*

No	D	H	W	W/H	U
SAM-PCA3496	51	19	22	1,16	19 (37)

*Discussion*

*Douvilleiceras variable* Tavani seems to be based upon the inner whorls of a macroconch, showing the irregular ornament of the early growth stages, with 1–2 weak ribs between main ribs, and the more regular ornament of the macroconch on the adoral half of the last whorl. Tavani's (1949) species is undoubtedly very close to *D. mammillatum* (Schlotheim), and D'Orbigny's (1841, pl. 72 (fig. 5)) *Ammonites mammillaris* seems to be a typical example of this species. Indeed, even in the present collection, several individuals seem to combine the characters of *D. variable* and *D. mammillatum aequinodum* (Fig. 15E). This tends to suggest that *D. variable* is merely another morphotype within a polytypic *D. mammillatum*. While this is the writer's belief, opposition to the gross 'lumping' that such synonymies would involve from one of the referee's of this paper has led to the adoption of a somewhat more cautious approach. *Douvilleiceras variable* is here maintained as a distinct species because the writer does not believe in sympatric subspecies.

*Douvilleiceras subleightonense* Mirzoyev is a gigantic macroconch attaining some 600 mm diameter, whose inner whorls show the irregular ribbing typical of this species, and it may reasonably be included here. The *Douvilleiceras* sp. B of McLearn (1972, pl. 13 (fig. 1), pl. 26 (fig. 1)) is another macroconch that may belong here. Yet another macroconch whose inner whorls are reminiscent of this species is *D. pseudinaequinodum* Collignon. *Douvilleiceras ivernoisi* (Coquand) (Pictet & Renevier, 1854, pl. 2 (fig. 1)) is somewhat similar to *D. variable*, but lacks the pronounced lateral tubercle.

*Douvilleiceras inaequinodum* (Quenstedt, 1849)

Figs 4C–D, F, 6A, 7A–C, 9D, F, H, 10E, 11A, 12C–D, G–H

*Ammonites mammillaris* Schlotheim, d'Orbigny, 1841: 249, pl. 73 (figs 1–3 only).

*Ammonites monile inaequinodum* Quenstedt, 1849: 138, pl. 10 (fig. 1).

*Douvilleiceras inaequinodum* (Quenstedt) Parona & Bonarelli, 1897: 95, pl. 4 (fig. 6), pl. 13 (fig. 6). Collignon, 1950: 46, fig. 2; 1963: 114, pl. 285 (fig. 1245). Casey, 1962: 282, fig. 95c.

*Douvilleiceras orbignyi* Hyatt, 1903: 110. Casey, 1962: 279, pl. 40 (figs 6–8), pl. 42 (figs. 12–13). Howarth, 1965: 345, pl. 1 (fig. 5).

*Douvilleiceras mammillatum* var. *baylei* Spath, 1923: 70, pl. 5 (fig. 4).

*Douvilleiceras* aff. *inaequinodum* (Quenstedt) Spath, 1923: 73, pl. 4 (fig. 5).

? *Douvilleiceras* cf. *inaequinodum* (Quenstedt) Scott, 1937: 35, pl. 6 (figs 3–4), fig. 1.

*Douvilleiceras alternans* Casey, 1962: 282, pl. 42 (fig. 1), fig. 1021.

? *Douvilleiceras magnodosum* Casey, 1962: 284, pl. 42 (fig. 4), fig. 102k.

? *Douvilleiceras charshangense* Mirzoyev, 1967: 54, pl. 7 (figs 1–5).

? *Douvilleiceras* sp. A, McLearn, 1972: 67, pl. 12 (fig. 1).

*Douvilleiceras* cf. *orbignyi* Hyatt, Young, 1974: 188, pl. 5 (figs 9–10).

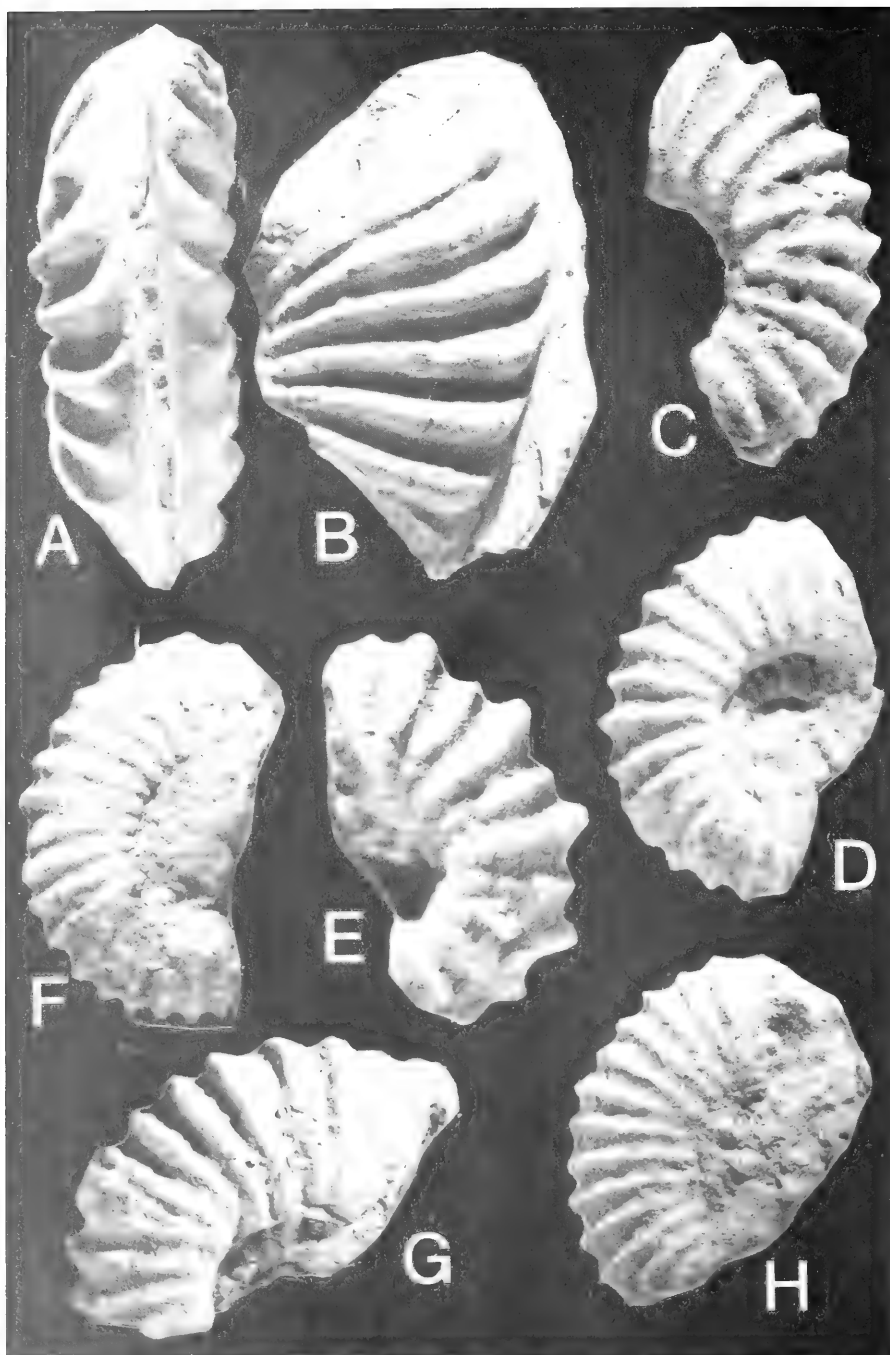


Fig. 10 A-B. *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White), SAM-PCA2656, a *douglasi* morphotype which very closely approaches *O. mirapelianum* (d'Orbigny),  $\times 0.75$ . C-D. F-H. *Douvilleiceras* *mammillatum aequinodum* (Quenstedt). C. SAM-PCA3494. D. SAM-PCA3503. F. SAM-PCA3527. G. SAM-PCA3499. H. SAM-PCA3518. E. *Douvilleiceras inaequinodum* (Quenstedt), SAM-PCA3504. All  $\times 1$  unless otherwise stated.

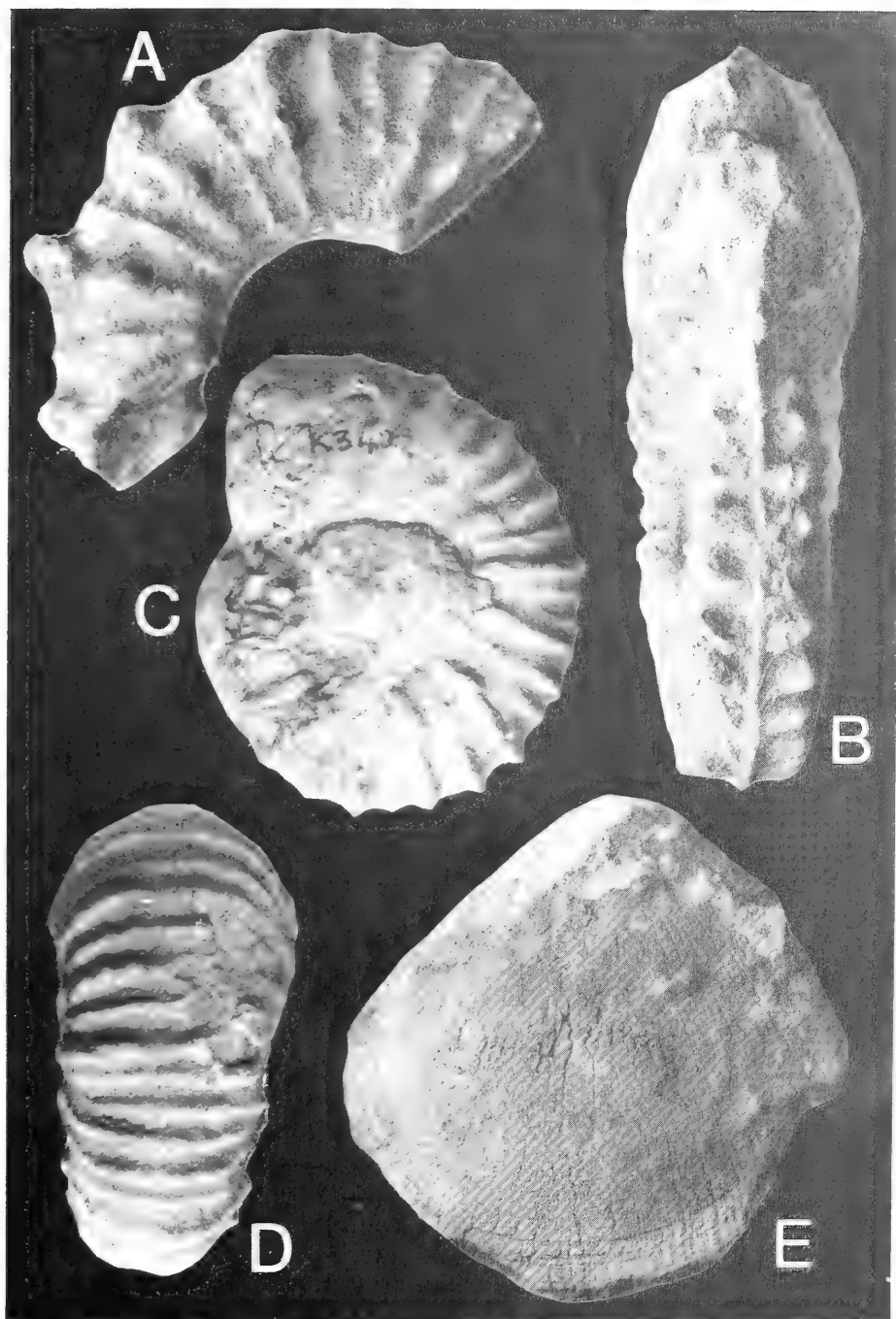


Fig. 11. A. *Douvilleiceras inaequinodum* (Quenstedt), SAM-PCA3458, a specimen showing the increased number of intercalatories in maturity. B. *Oxytropidoceras* (*Mirapelia*) *sergipense* (White). Front view of SAM-PCA5444. C-D. *Douvilleiceras variabile* Tavani, SAM-PCA3422. E. *Plagiostoma* sp., SAM-PCA5490, from the *Oxytropidoceras* assemblage. All  $\times 1$ .

*Material*

Eighteen specimens, SAM-PCA3435–3438, 3444, 3451, 3453, 3456–3458, 3466, 3492, 3495, 3498, 3504, 3512, 3525, and 3535, all preserved as limestone steinkerns.

*Description*

This is a rather common species in the Angolan fauna. The shell is generally moderately evolute, inflated, with a subreniform, depressed whorl section. The umbilicus is rather wide, with steep umbilical walls and well-rounded shoulders. Ornament comprises sparse, distant, slightly rursiradiate ribs, which are narrower than the interspaces and alternate weak and strong. The strong main ribs are ornamented with small umbilical tubercles, a prominent lateral tubercle, and exaggerated ventral bulges, each with three spiral notches. In juveniles, only two spiral notches are present on the ventral bulges. With ontogeny, a small tubercle appears between the umbilical and mid-lateral tubercle and another between the latter and the ventrolateral bulges. Thus, there may be up to eight crenulations per rib in adults. The fine intercalated ribs, usually one and rarely two, between main ribs generally lack discernible ornament (on the weathered internal moulds), but in some individuals develop a small but distinct mid-lateral tubercle. In the early growth stages (Figs 4C–D, 12G–H) the intercalated ribs may be absent or so fine that there is difficulty in distinguishing this species from *D. mammillatum* s.s. There are probably about 7–8 main ribs per half whorl. The ventral sulcus is moderately broad and rather deep.

*Discussion*

Casey (1962) considered *D. inaequinodum* to be a typical Middle Albian species, distinguished from the earlier *D. orbignyi* Hyatt and *D. alternans* Casey in having two, not one, fine intercalatories between main ribs. However, material from the *Isohoplites eodentatus* nodule bed at Bully, Pays de Bray, France (Fig. 12C–D), shows that at least some individuals of *D. inaequinodum* have but a single fine intercalated rib between main ribs. In view of this, and because Casey (1962) has noted transitions between *D. alternans* and *D. orbignyi*, the latter two species are included in the synonymy of *D. inaequinodum*. Until the population structures of the *mammillatum* Zone assemblages are better known, it is perhaps prudent to retain *orbignyi* as a subspecies for these forms, as *D. inaequinodum orbignyi* Hyatt.

Family **Anisoceratidae** Hyatt, 1900

Genus *Protanisoceras* Spath, 1923

Type species *Hamites raulinianus* d'Orbigny, 1842; by original designation  
*Protanisoceras* sp.

Fig. 13B

*Material*

A single fragment, SAM-PCA3461, preserved as a limestone steinkern from the *Douvilleiceras* assemblage.



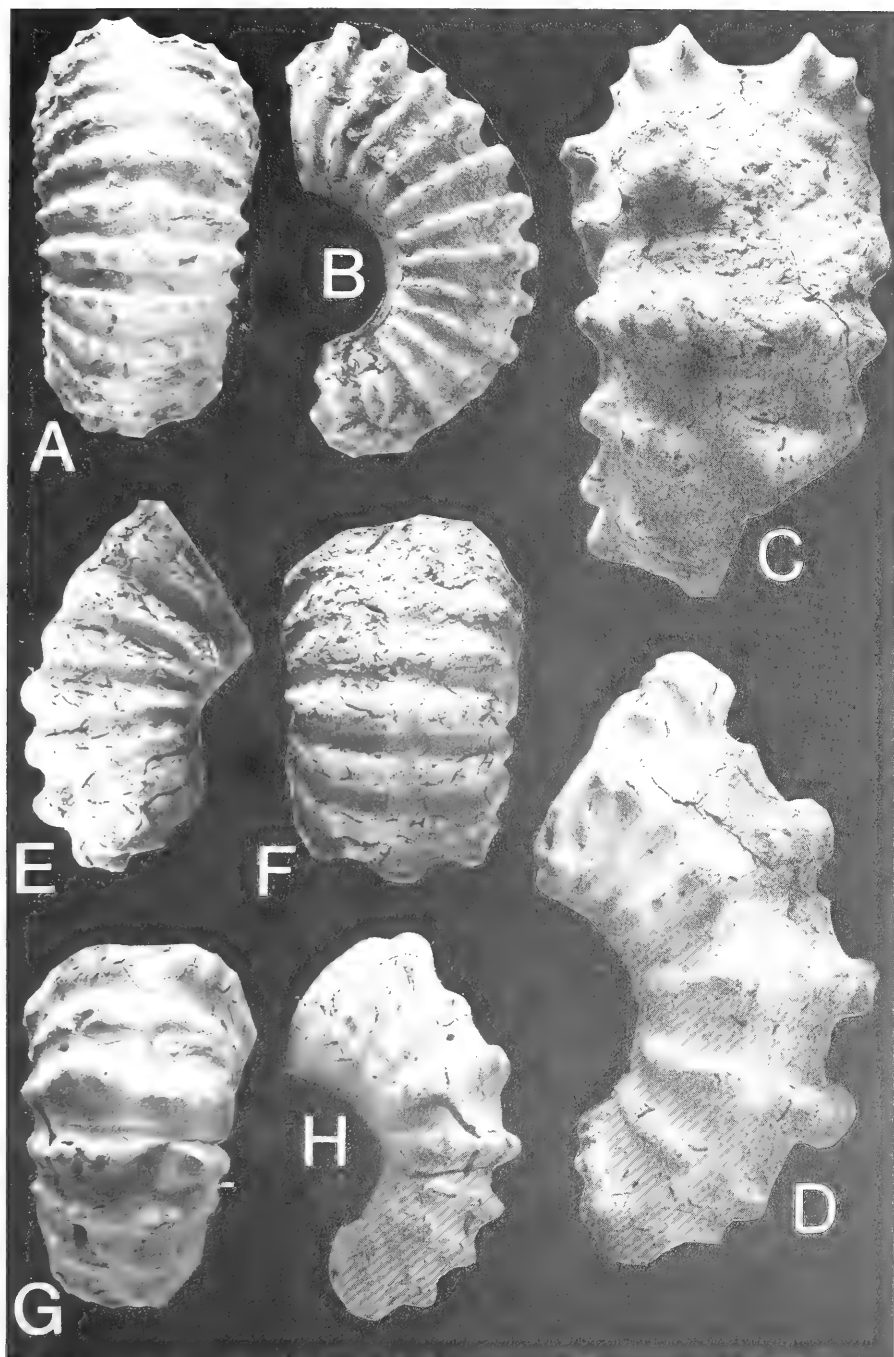


Fig. 12. *Douvilleiceras* spp. from the *I. eodentatus* Nodule Bed, Bully, Pays de Bray, France (Bed e of Destombes *et al.* 1977). A-B. *Douvilleiceras mammillatum aequinodum* (Quenstedt), OUM-KZ976. C-D, G-H. *Douvilleiceras inaequinodum* (Quenstedt). C-D. OUM-KZ982. G-H. OUM-KZ981. E-F. *Douvilleiceras mammillatum mammillatum* (Schlotheim), OUM-KZ979. All  $\times 1$ .

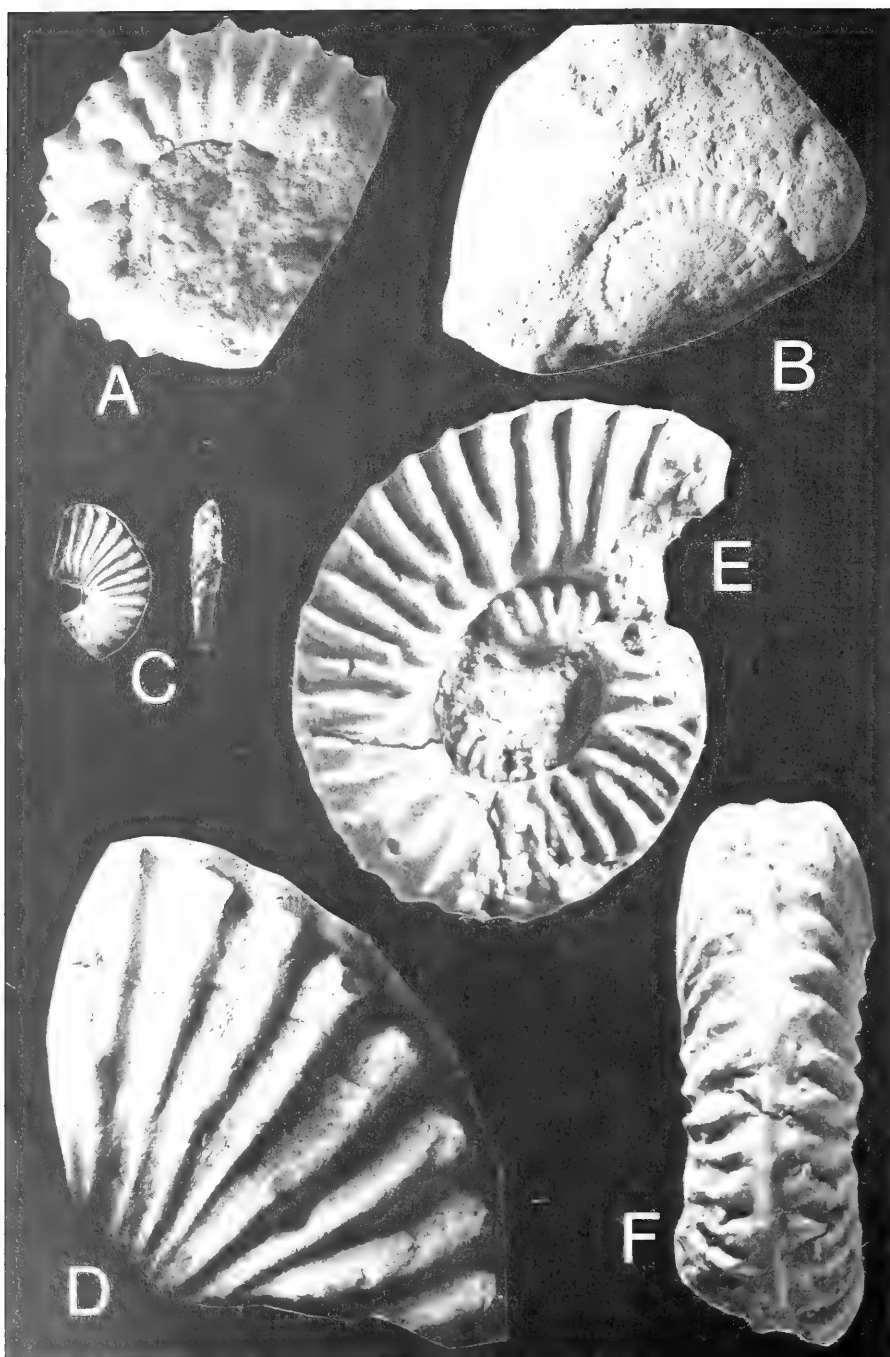


Fig. 13. A. *Douvilleiceras variable* Tavani, SAM-PCA3472. B. *Protanisoceras* sp., SAM-PCA3461. C-D. *Venezoliceras acostae* (d'Orbigny), syntypes in the D'Orbigny Collection, NHMP. E-F. *Mortoniceratoides rigidus* (Spath), the holotype, BM-C34879. All  $\times 1$ .

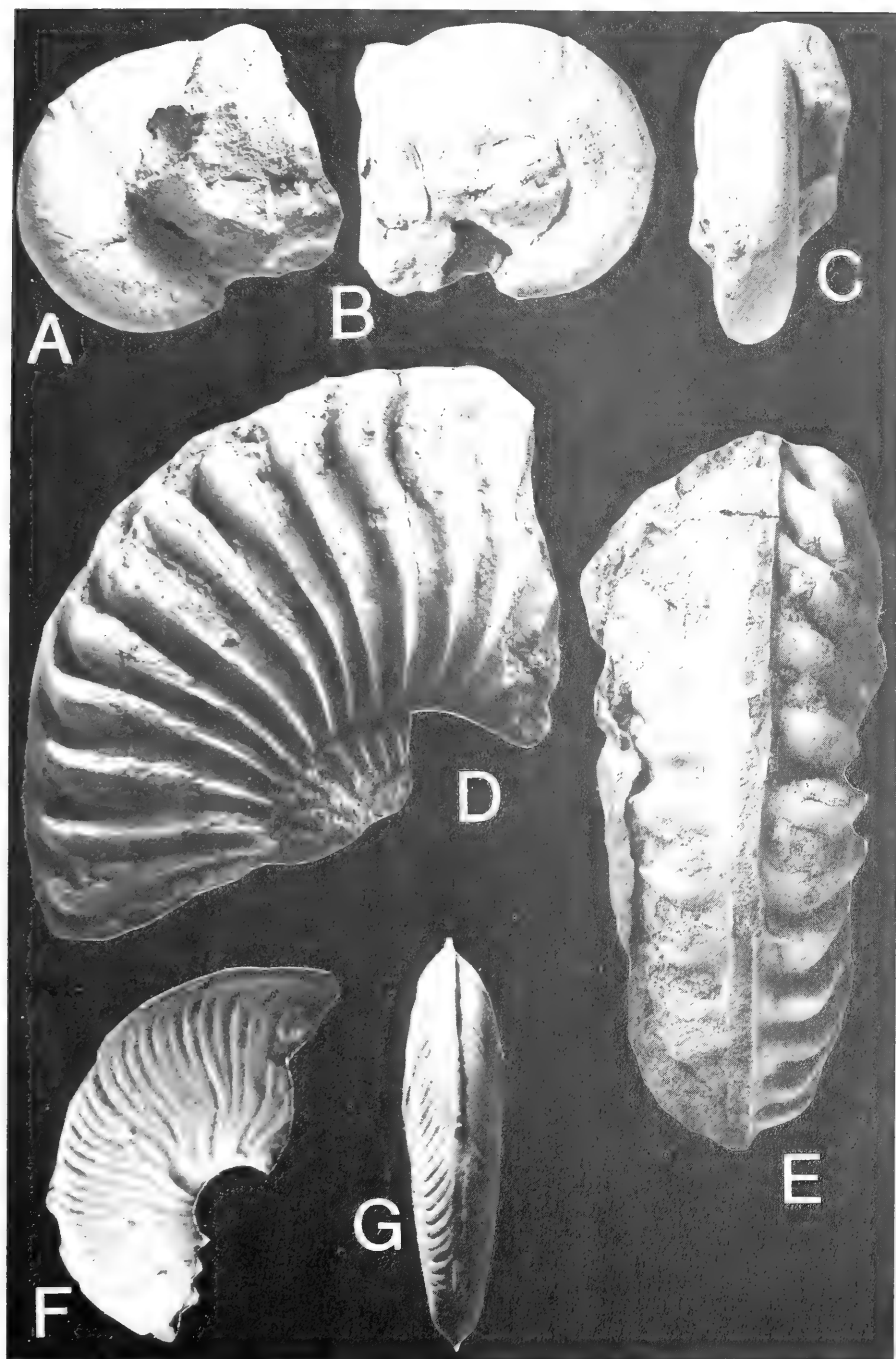


Fig. 14. A–C. *Puzosia bistricta* (White), the lectotype, designated herein, USNMNH 12102, from Porto dos Barcos, Mariom, Sergipe, Brazil. D–E. *Oxytropidoceras* (*Mirapelia*) *mirapelianum* (d'Orbigny), NHMP-5758, in the D'Orbigny Collection, from Clar, France. F–G. *Oxytropidoceras* (*Oxytropidoceras*) *carbonarium* (Gabb), the holotype of *O. cantianum* Spath, BM-C12501, from the *D. cristatum* Zone at Folkestone.

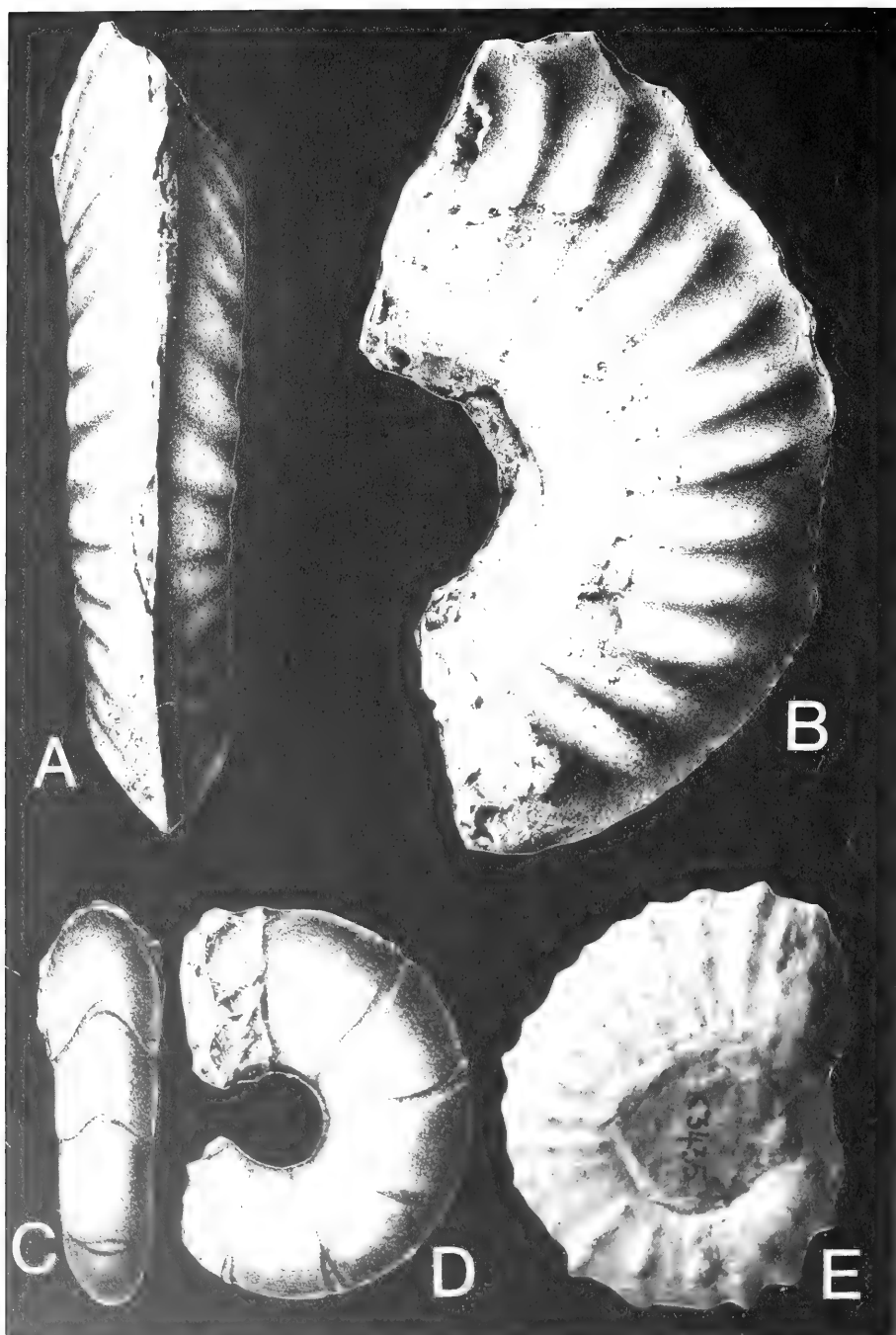


Fig. 15. A–B. *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White), the lectotype, designated herein (after White 1887). C–D. *Puzosia bistricta* (White), copy of White's (1887) original illustration. E. *Douvilleiceras* cf. *variabile* Tavani, SAM-PCA3435. Note the abrupt change from a *variabile* to an *aequinodum*-type ornament. All  $\times 1$ .

### Description

The short curved fragment gives the impression of having had a compressed, elliptical whorl section. Ornament comprises rather broad, low, rursi-radiate ribs, wider than the interspaces, each bearing a distinct ventrolateral tubercle. There are about three ribs in a distance equal to the whorl height.

### Discussion

The available specimen is too fragmentary to warrant comparison with the many described species of *Protanisoceras*, but does provide the first record of this genus from Angola.

Family **Desmoceratidae** Zittel, 1895

Subfamily Puzosiinae Spath, 1922

Genus *Puzosia* Bayle, 1878

Type species *Ammonites mayoriana* d'Orbigny, 1842  
(= replacement name for *A. planulatus* J. de C. Sowerby  
non Schlotheim 1820 nec Schuebler 1830);  
by subsequent designation

*Puzosia bistricta* (White, 1887)

Figs 2A–B, 3C–D, 14A–C, 15C–D, 16, 17B–C, 18C–D, 19A–B

*Ammonites bistrictus* White, 1887: 216, pl. 23 (figs 5–8).

*Ammonites hopkinsi* Forbes, White, 1887: 213, pl. 21 (figs 1–3), pl. 22 (fig. 5).

*Puzosia welwitschia* Choffat (in Choffat & de Loriol), 1888: 68, pl. 2 (fig. 4).

*Puzosia garajauana* Maury, 1930: 289; 1936: 234, pl. 20 (figs 1–3). Beurlen, 1970: 446, pl. 1 (figs 1–3), figs 1–2.

*Puzosia rosarica* Maury, 1936: 236, pl. 26 (figs 2–3). Beurlen, 1970: 446, pl. 1 (figs 4–5), figs 3–6.

### Material

Thirty specimens, SAM-PCA2623, 2625–2626, 2636, 2638, 2649, 2652, 2660–2661, 3412, 3415–3416, 3419–3421, 3424, 3426, 3429, 3432, 3434, 3449, 3452, 3474–3475, 3635, 3749, 3751, 3753, 3757, all preserved as limestone steinkerns and mostly from the beds with *Douvilleiceras*.

### Description

Shell moderately large, compressed, with a moderately narrow umbilicus (21–27% of the diameter). The umbilical walls are steep, with evenly rounded shoulders, and the flanks slightly convex, almost flat. Maximum width is just below mid-flank and the whorl section (Fig. 16) is elliptical. There are generally about 7–8 straight, prorsiradiate constrictions per whorl, which flex forward just before crossing the venter. Characteristically, the whorls show an increase in inflation immediately after a constriction. The constrictions are broad and shallow, typically with a peculiar wedge-shaped rib intercalated across the

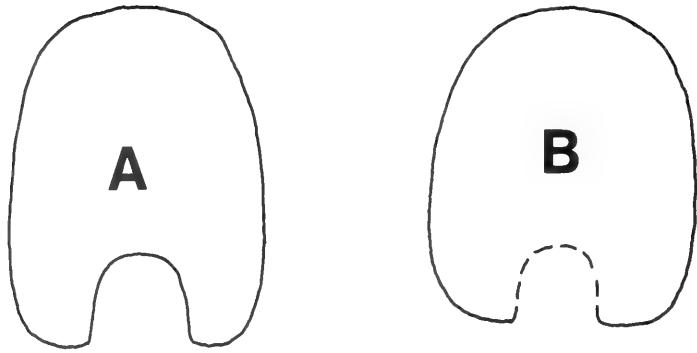


Fig. 16. Whorl sections of *Puzosia bistricta* (White). A. SAM-PCA2626. B. SAM-PCA3412. Both  $\times 1$ .

venter in the middle growth stages. Because of the nature of preservation, the true characters of the ribbing are difficult to assess. They first appear on the outer parts of the flanks at about 60–80 mm diameter, and coarsen and become more prominent across the venter with age. On SAM-PCA2623 there are twenty fine ribs between successive constrictions whereas on SAM-PCA2649 there are only seven.

#### Discussion

*Puzosia welwitschia* Choffat is clearly conspecific with the *Ammonites hopkinsi* figured by White (1887), which was later renamed *Puzosia garajauana* by Maury (1930). They almost certainly merely represent the middle and adult growth stages, respectively, of *P. bistricta* (White) (Fig. 14A–C). *Puzosia rosarica* Maury occurs with *P. bistricta* and is based upon juvenile material which probably does not warrant specific separation. There are few species of *Puzosia* with which this highly distinctive Middle Albian species may be confused.

Family **Brancoceratidae** Spath, 1933

Subfamily Mojsisovicziinae Hyatt, 1903

Genus *Oxytropidoceras* Stieler, 1920

Type species *Ammonites roissyanus* d'Orbigny, 1841;

by original designation

#### Discussion

The most comprehensive treatment of this genus is that of Young (1966). He included *Androiavites* Collignon (1936) in the synonymy of *Adkinsites* and provided the following emended diagnosis of *Manuaniceras*: 'Ammonites with

few whorls, which are much higher than wide; there are many flat ribs at some stage in the ontogeny, and ribs may be single or bifurcating at one or more positions at the umbilicus or on the flank' (p. 95).

Unfortunately, however, flat-topped ribs are not a feature of *Manuaniceras manuanense* (Spath) (Fig. 31B) and hence cannot be used in the formulation of a diagnosis of this taxon. As Young (1966) rightly pointed out, *Manuaniceras* is a direct descendant of the *Oxytropidoceras roissyanum* group, which mainly differs in showing rib bifurcation at various levels on the flank. The differences are slight and the writer finds little use for the genus/subgenus *Manuaniceras* which is here included in the synonymy of *Oxytropidoceras* s.s.

As noted by Young (1966), and is clear both from the stratigraphic record and described species of *Oxytropidoceras*, the history of this taxon is one of parallel evolution between two species groups. The *O. mirapelianum* (d'Orbigny) (Fig. 14D-E herein) plexus comprises *O. mirapelianum* (d'Orbigny), *O. evansi* (Spath), *O. douglasi* Knetchel, *O. buarquianum* (White), *O. sergipense* (White), *O. packardi* Anderson, *O. boulei* Collignon, *O. paucituberculatum* (Collignon), and *O. colcanapi* Collignon. These are all coarsely ribbed species without bifurcation or intercalation of the simple ribs. To this group the writer would assign the strongly derived species of *O. salasi* Young, *O. stenzeli* Young, and *O. pandalense* Young in which the adult whorls become densicostate, with occasional bifurcation and intercalation, and thus homoeomorph 'Manuaniceras'. The other species group is that of *O. roissyanum* (d'Orbigny) in which the ribs are fine, with frequent bifurcation at or above the umbilical edge, and which includes all the described 'Manuaniceras' species of Young (1966). Since these two plexi appear almost simultaneously at the base of the Middle Albian (Owen 1971) and evolved in parallel until their final extinction in the early late Albian, the writer believes they merit subgeneric distinction. The writer proposes, therefore, the new subgenus *Oxytropidoceras* (*Mirapelia*) to include all those coarse-ribbed species assigned to the *mirapelianum* plexus above, with *Ammonites mirapelianus* d'Orbigny as type species. *Oxytropidoceras* (*Mirapelia*) *sergipense* (White) provides the root-stock from which the typical coarse-ribbed species of *Venezoliceras* (of which *Tarfayites* Collignon (1966) is a synonym), such as *V. venezolanum* (Stieler), *V. acostae* (d'Orbigny) (Fig. 13C-D), *V. heueri* Young, *V. obscurum* Young, *V. texanum* Young, *V.?* *chihuahuaense* (Böse), *V. kiowanum* (Twenhofel), *V. umsinenense* (Van Hoepen), *V. bituberculatum* (Collignon), *V. madagascariense* (Collignon), *V. robustum* (Renz) and *V. clavicostatum* (Renz), are derived. However, as noted by Young (1966) and Renz (1968), there is a second group of *Venezoliceras*, typified by *Venezoliceras harrisoni* Benavides-Cáceres, in which the early and middle growth stages show dense, bifurcating, nontuberculate ribbing, with a flank tubercle developed only at relatively large diameters. This group, which also includes *V. acutocarinatum* (Shumard), *V. multicostatum* (Renz), *V. intermedium* (Renz), and *V. karsteni* (Stieler), seems to have its ancestry in 'Manuaniceras' and, as such, its similar-

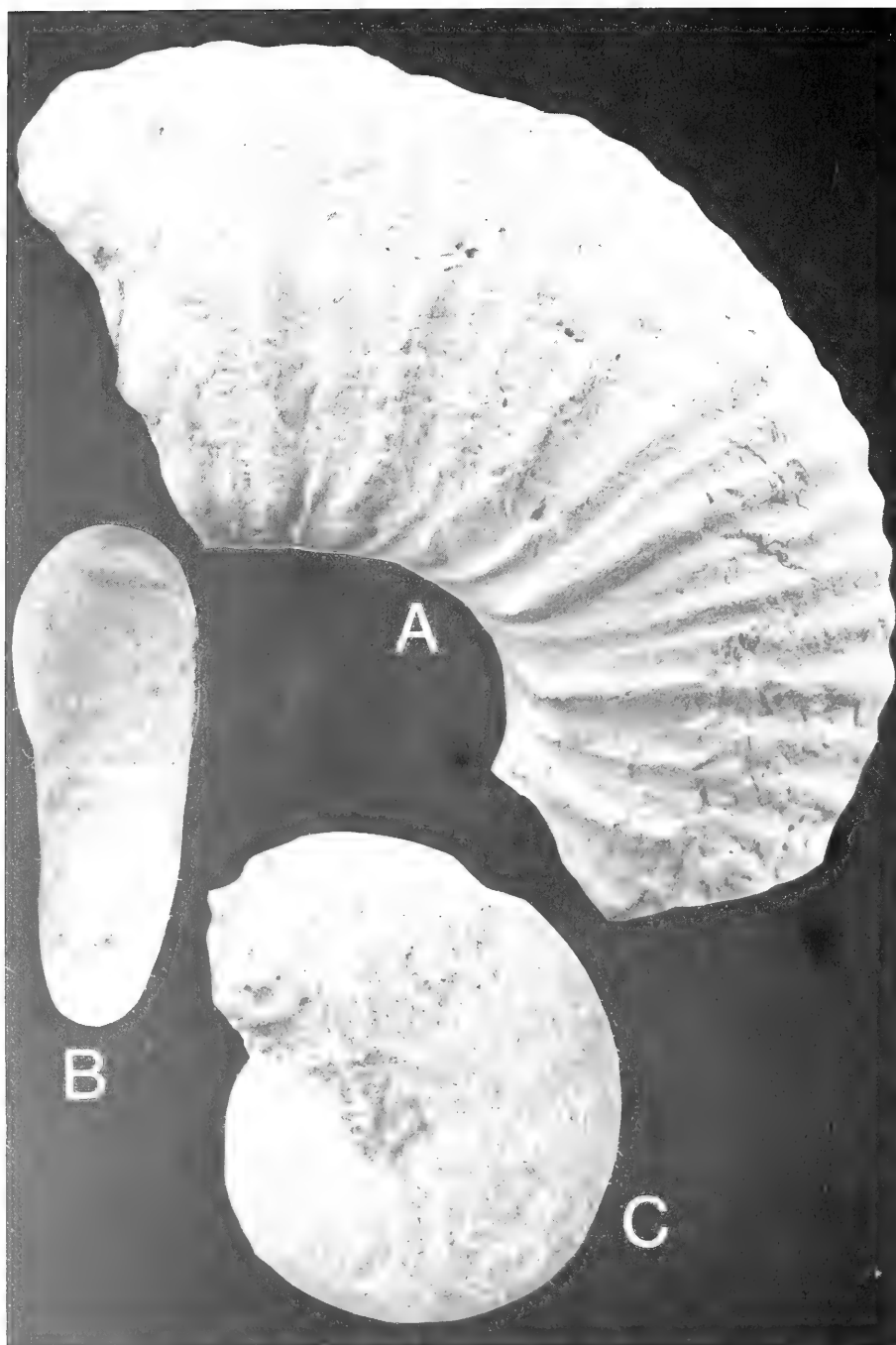


Fig. 17. A. *Douvilleiceras mammillatum aequinodum* (Quenstedt), SAM-PCA3423, a macroconch fragment,  $\times 0.67$ . B-C. *Puzosia bistricta* (White), SAM-PCA5447,  $\times 1$ .



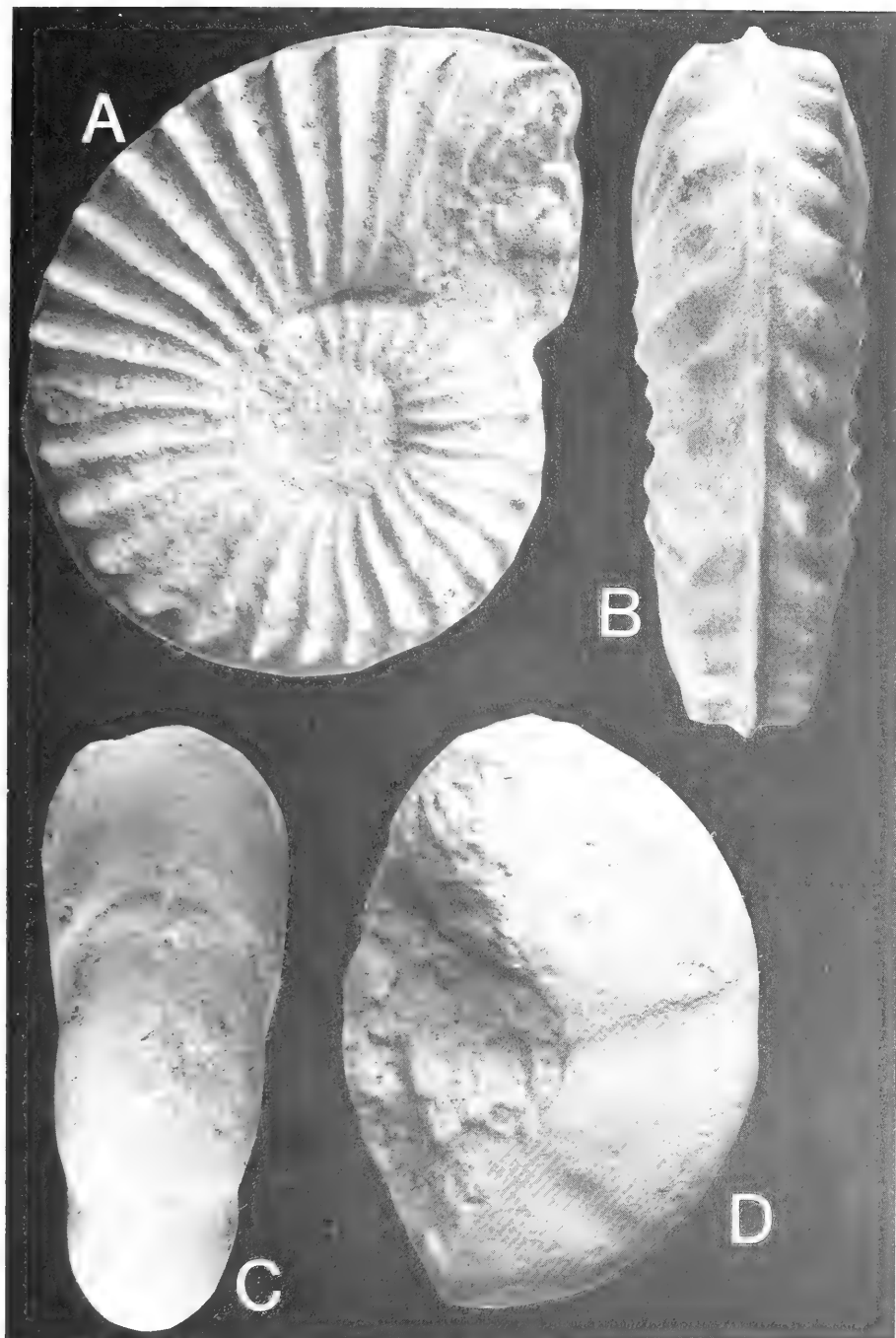


Fig. 18. A-B. *Oxytropidoceras (Mirapelia) sergipense* (White), SAM-PCA5444. C-D. *Puzosia bistricta* (White), SAM-PCA3420. Both  $\times 1$ .

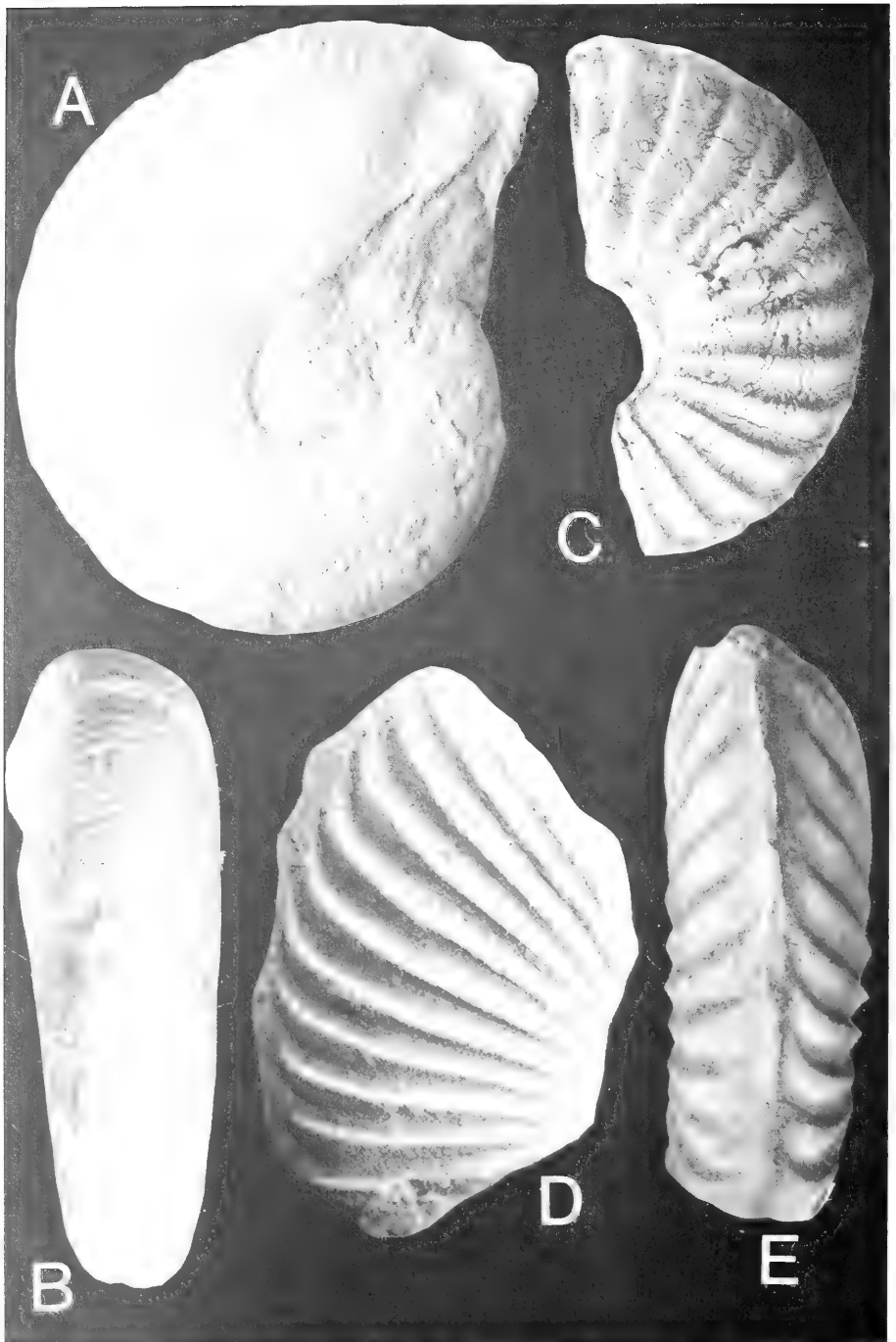


Fig. 19. A-B. *Puzosia bistricta* (White), a specimen in the South African Museum. C-E. *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White). C. SAM-PCA5446, a *douglasi* morphotype. D-E. SAM-PCA3603, a *salasi* morphotype. All  $\times 1$ .

ity to the *V. acostae* group is due to convergence. Consequently, the writer proposes the new subgenus *Oxytropidoceras* (*Benavidesites*), with type species *Venezoliceras harrisoni* Benavides-Cáceres (1956: 460, pl. 53 (fig. 6)), for this homoeomorphic development.

The origins of *Adkinsites* are obscure; Young (1966) would derive this taxon from *Dipoloceras*, but this view is unacceptable to the writer. They could represent derivatives of the more densely ribbed *O. (Mirapelia)* species or of paucicostate *Oxytropidoceras* s.s., or of both. This taxon is not present in the Angolan material and hence the writer defers judgement. Because of its obscure, and perhaps polyphyletic origins, Young (1966) is followed in treating *Adkinsites* as a genus, of which *Androiavites* Collignon (1936) is a synonym.

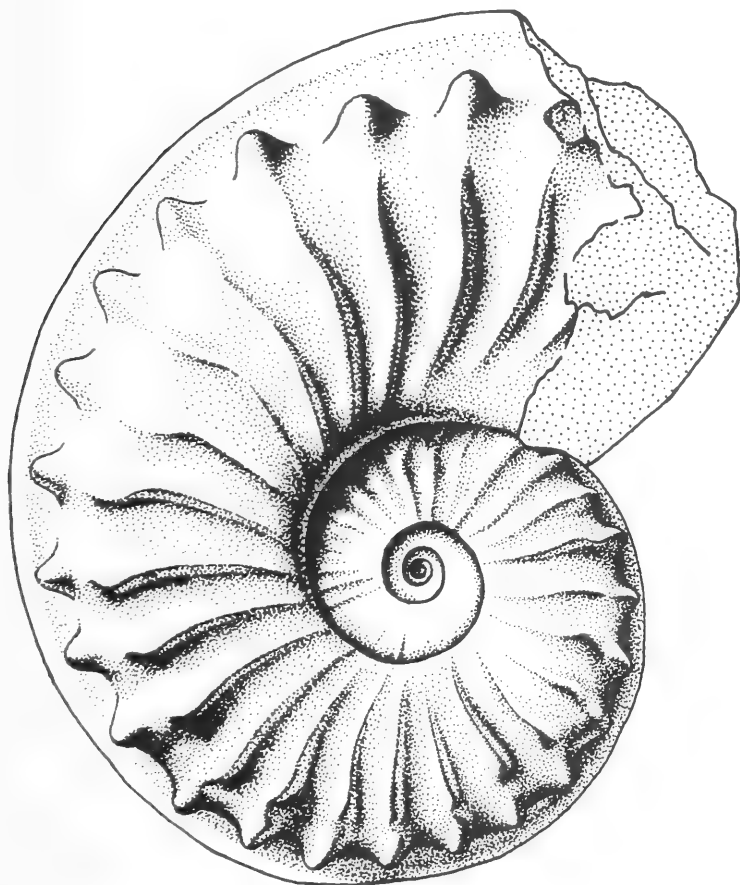


Fig. 20. *Mojsisoviczia* cf. *ventanillensis* (Gabb). A reconstruction based upon BM-C78865, from the Middle Albian of Zululand.  $\times 1$ .

*Mortoniceras* (*Pervinqueria*) *rigidum* Spath (1933: 413, figs 142, 144f) (Fig. 13E–F herein) is recorded from Bed VIII (= *D. cristatum* Zone) at Folkestone and is, as such, the earliest described species of *Mortoniceras*. It differs from the latter genus, however, in having rather fine, distant ribs with a steep adoral surface and a sloping adapical surface, lacking any sign of a mid-lateral tubercle, and in lacking the doubling of the ventrolateral tubercle so typical of *Mortoniceras*. Indeed, the characters of ornament ally *M. rigidum* with the *Mojsisovicziinae* and it seems likely that *M. rigidum* is closely allied to, and perhaps descended from *Adkinsites*. Since the origins of *Mortoniceras* are generally thought to lie in *Dipoloceras* (Wright in Arkell *et al.* 1957), the similarities are due to convergence. It is proposed, therefore, to make *M. rigidum* Spath the type species for the new genus *Mortoniceratoides*. It differs from *Adkinsites* in having a subquadrate whorl section, low ventral keel, and with a symmetrically bifid first lateral saddle.

As discussed by Kennedy & Cooper (1977), *Mojsisoviczia* and *Falloticerias* are closely allied and, on the basis of the available evidence, *Falloticerias* may reasonably be considered a neotonous offshoot of *Mojsisoviczia*. From their first appearance (Fig. 20), the ribs of *Mojsisoviczia* are coarse and simple and the origins of this taxon seem to lie in *O. (Mirapelia) sergipense* (White) or a closely allied form.

*Dipoloceras* is believed by Spath (1931) to have descended from *Oxytropidoceras*, and is included in the *Mojsisovicziinae* by Wright (in Arkell *et al.* 1957). However, the inflated whorls, low ventral keel and strong spiral ornament suggest a closer relationship with the *Mortoniceratinae*, to which subfamily *Dipoloceras* is here referred. The suggested phylogeny within the *Mojsisovicziinae*, as here interpreted, is shown as Figure 21.

### *Oxytropidoceras (Mirapelia) buarquianum* (White, 1887)

Figs 3A–B, 6B, 10A–B, 15A–B, 19C–E, 22D–E, 23A–D, F–G, 24–25, 27

*Ammonites buarquianus* White, 1887: 222, pl. 24 (figs 3–4 only), pl. 25 (figs 7–8).

*Schloenbachia* cf. *roissiana* (d'Orbigny) Douvillé, 1906: 147, pl. 4 (fig. 3).

*Schloenbachia* aff. *belknapi* (Marcou) Douvillé, 1906: 148, pl. 2 (fig. 4).

*Schloenbachia* cf. *belknapi* Böse (non Marcou) Douglas, 1921: 269, pl. 17 (figs 1–2).

*Schloenbachia* cf. *chihuahuaensis* Böse, Douglas, 1921: 269, pl. 17 (fig. 3).

*Oxytropidoceras buarquianum* (White) Maury, 1936, pl. 19 (figs 2–3), pl. 25 (figs 5–6). Oliveira & Leonardos, 1943, pl. 27 (fig. 7). Young, 1966: 89, pl. 38 (figs 1–3). Beurlen, 1970: 464, pl. 5 (figs 1–2), fig. 18.

? *Oxytropidoceras colcanapi* Collignon, 1936: 182, pl. 18 (fig. 7), fig. 12k.

? *Oxytropidoceras* sp. juv., Spath, 1930: 61, pl. 9 (fig. 18).

*Oxytropidoceras packardi* Anderson, 1938: 198, pl. 50 (fig. 1).

*Oxytropidoceras douglasi* Knetchel, 1947: 106, pl. 24 (figs 1–4), pl. 25 (figs 1–2), pl. 26, pl. 27 (figs 2–3), pl. 28 (fig. 1). Young, 1966: 88, pl. 10 (figs 1, 5), pl. 36 (figs 3–4), pl. 37 (figs 4–5).

? *Oxytropidoceras involutum* Beurlen, 1952: 162, fig. 3.

? *Oxytropidoceras (Manuaniceras) involutum* Beurlen, 1970: 465, pl. 5 (fig. 4).

*Oxytropidoceras* sp. A, Beurlen, 1970: 464, pl. 5 (fig. 3).

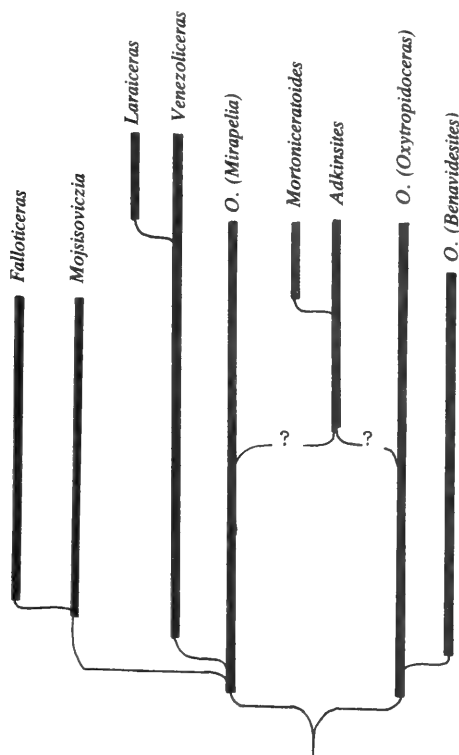


Fig. 21. A suggested phylogeny of the Mojsisoviciinae.

### Material

Forty-five specimens, including SAM-PCA2617–2618, 2621–2622, 2627–2630, 2632, 2637, 2640–2645, 2648, 2651, 2654, 2659, 2663, 2665–2667, 2669, 2673–2674, 2676, 2678, 2693, 3603–3604, 3755, 3762 and 3765, all preserved as limestone steinkerns.

### Description

The Angolan population is polytypic, with morphotypes agreeing exactly with *O. buarquianum* (White), *O. packardi* Anderson, *O. douglasi* Knetchel, *O. salasi* Young, and two specimens very closely approaching *O. mirapelianum* (d'Orbigny). The relationship between these morphotypes is shown in Figure 26. Because of this wide range of intraspecific variation, no single description will fully cover all morphotypes and hence they are described separately.

Two specimens, SAM-PCA2656–2657, very closely approach *O. mirapelianum* (d'Orbigny) and, with a typological approach, would be referred to that species. The Angolan examples (Fig. 10A–B) are, however, insensibly con-

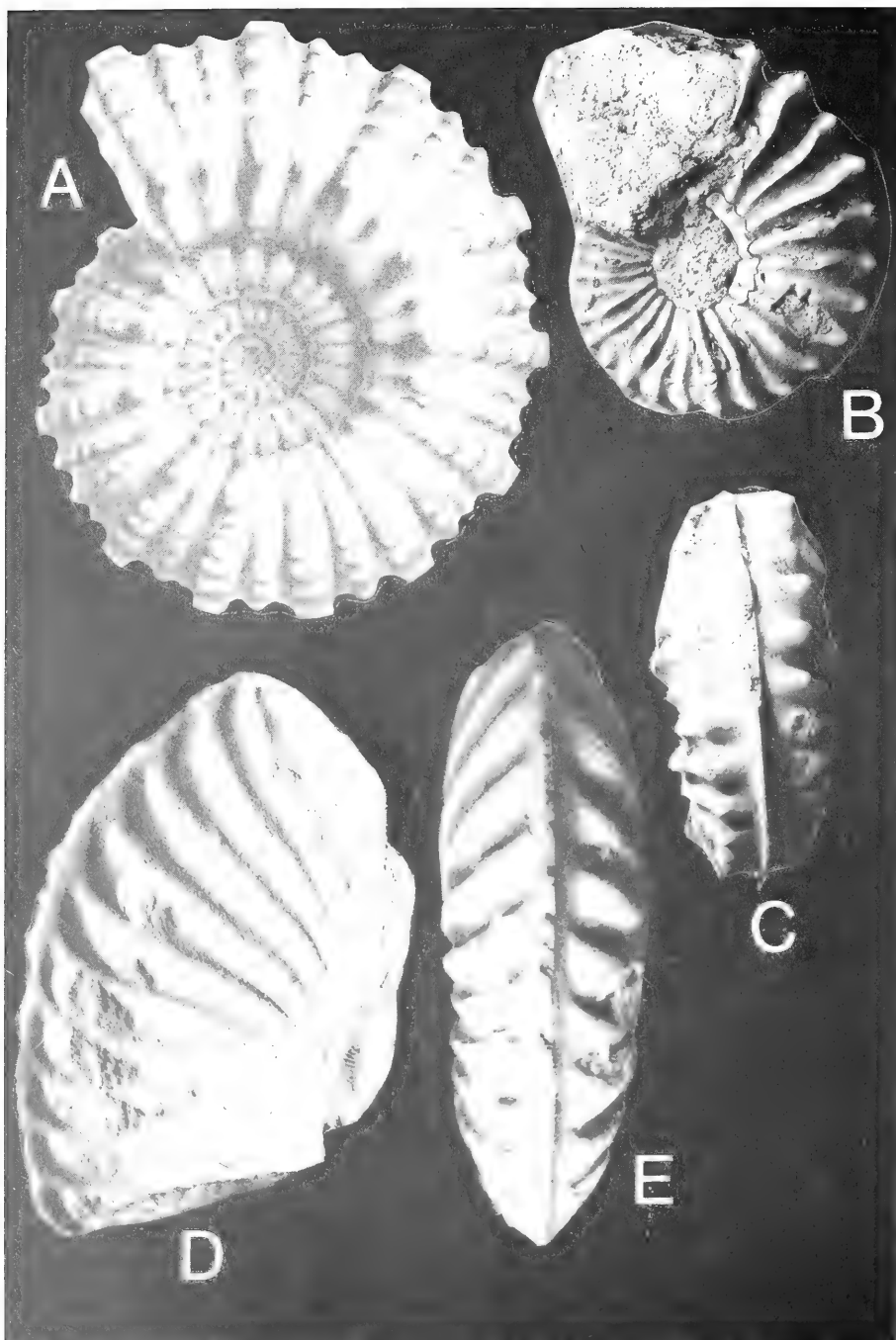


Fig. 22. A. *Douvilleiceras variable* Tavani, D'Orbigny's (1841, pl. 72 (fig. 5)) protograph,  $\times 0.5$ . B-C. *Mojsisoviczia delaruei* (d'Orbigny), NHMP-5761, one of D'Orbigny's syntypes from Clar, France. D-E. *Oxytropidoceras (Mirapelia) buarquianum* (White), SAM-PCA2641, a rather densely ribbed *douglasi* morphotype, thus transitional to *salasi* variants. All  $\times 1$  unless otherwise stated.

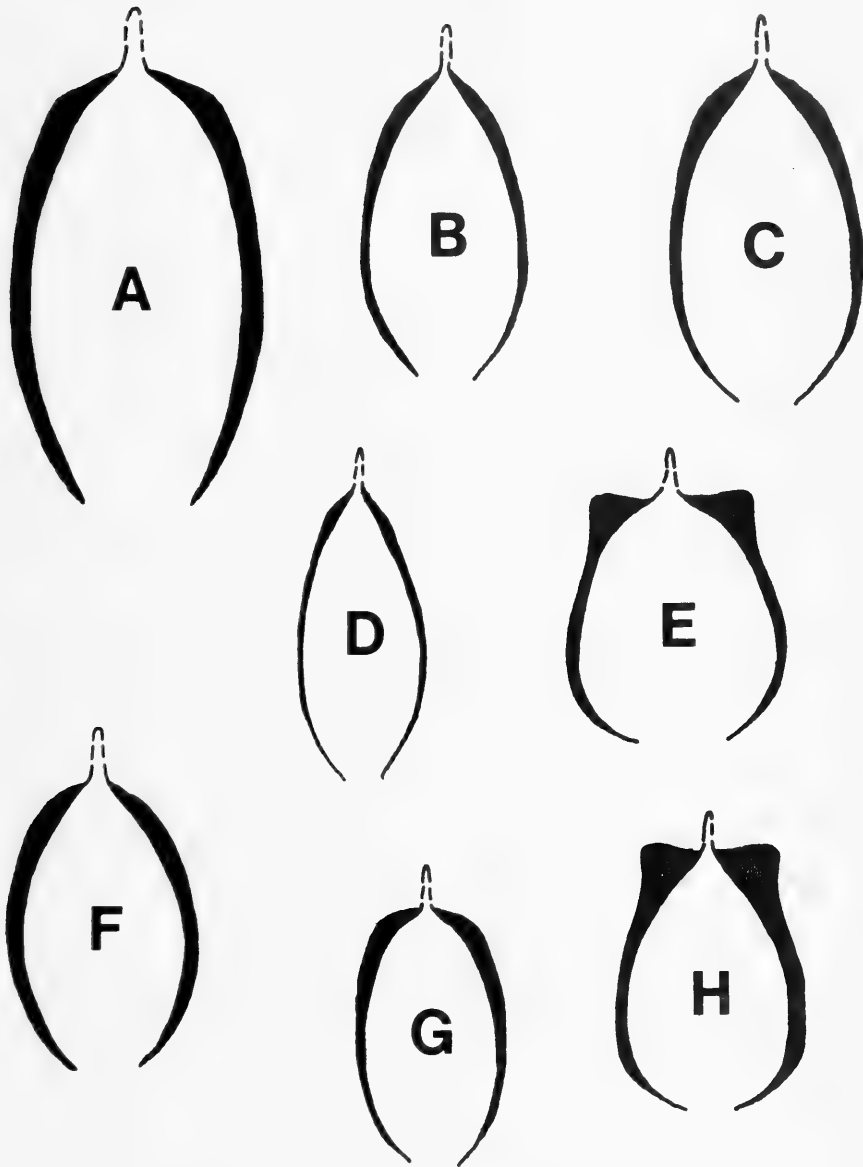


Fig. 23. A–D, F–G. Whorl sections of *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White). A. SAM-PCA2656, a *douglasi* morphotype. B. SAM-PCA2644, a *douglasi* morphotype. C. SAM-PCA3603, a *salasi* morphotype. D. SAM-PCA2693, a typical *buarquianum* morphotype. F. SAM-PCA2622, a *douglasi* morphotype. G. SAM-PCA2641, a *douglasi* morphotype. E, H. Whorl sections of *O. (M.) sergipense* (White). E. SAM-PCA5444. H. An unnumbered specimen in the South African Museum. All  $\times 1$ .



Fig. 24. *Oxytropidoceras (Mirapelia) buarquianum* (White), SAM-PCA3418, a *douglasi* morphotype.  $\times 1$ .



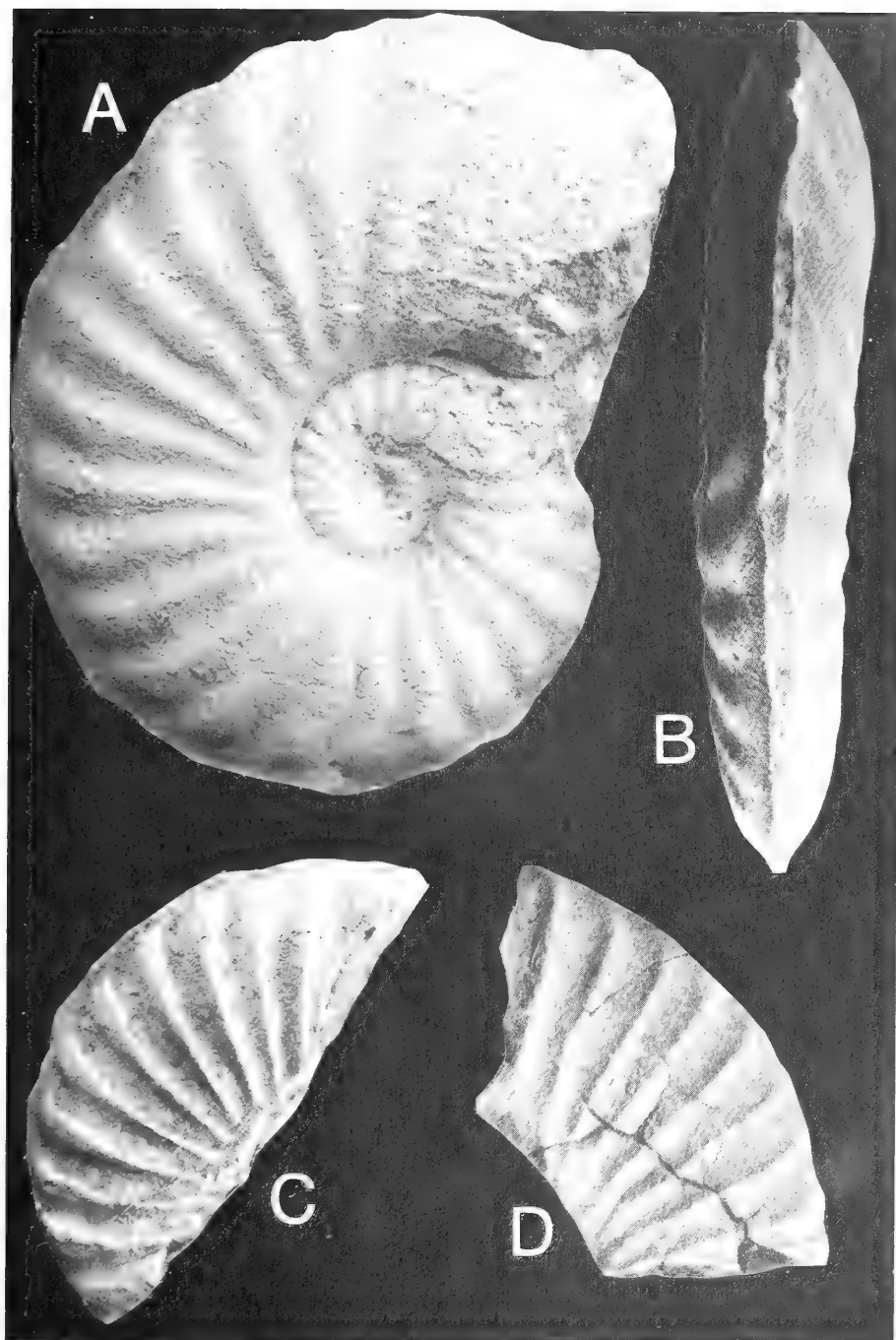


Fig. 25. *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White). A-B. SAM-PCA2693, the typical variant. C. SAM-PCA2674, a *douglasi* morphotype. D. SAM-PCA2665, a rather eroded, finely ribbed variant. All  $\times 1$ .

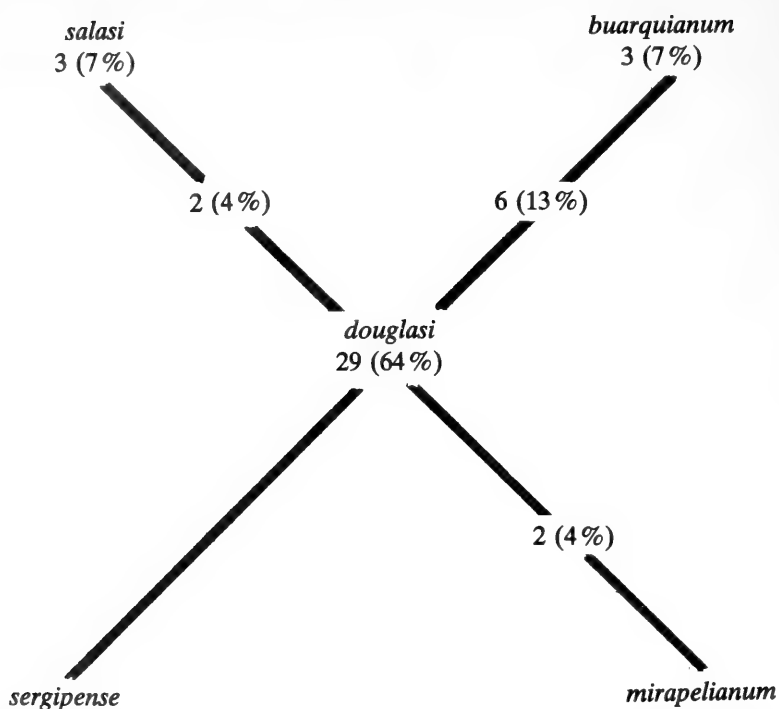


Fig. 26. Interrelationships between the *Oxytropidoceras* (*Mirapelia*) *buarquianum* morphotypes and closely allied species. Figures given are the number of morphotypes in the present collection followed, in parentheses, by the percentage which they constitute.

nected to *O. douglasi* morphotypes and differ from D'Orbigny's (1841) species (Fig. 14D–E) in being slightly less inflated and in having the ribs somewhat thicker at the umbilical margin.

The *O. douglasi* morphotypes are the most abundant in the present collection and show the following features: shell compressed, evolute (umbilicus 24–28% of the diameter), with a compressed, lanceolate whorl section and a prominent, sharp, high, siphonal keel. The umbilicus is rather wide and shallow, with fairly steep umbilical walls and evenly rounded shoulders at moderate diameters. The umbilical walls become increasingly inclined with growth and, in maturity, the umbilical shoulder is gently rounded and indistinct. At this stage, the outer whorl covers about 15 per cent of the ribbing of the preceding whorl. Maximum width is slightly below mid-flank in most individuals. Ornament comprises simple, generally flexuous ribs, which arise at the umbilical seam but only become prominent on the flanks. All ribs are slightly prorsiradate, steepest adorally and convex adapically, and broaden towards the ventrolateral shoulders where they swell but do not form tubercles.

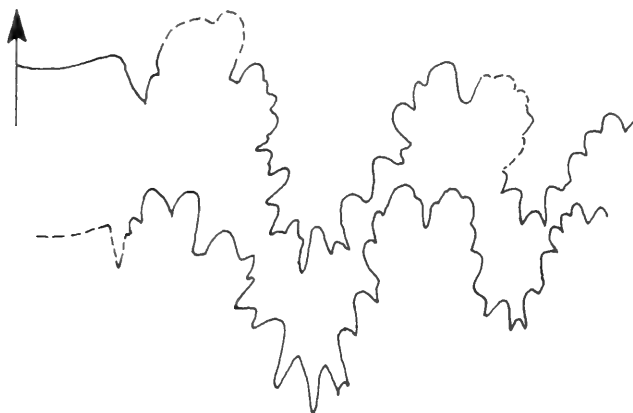


Fig. 27. *Oxytropidoceras* (*Mirapelia*) *buarquianum* (White). The suture line, somewhat simplified by erosion, of SAM-PCA2693, a *douglasi* morphotype. Approx.  $\times 2$ .

There are 13–20 ribs per half whorl. A somewhat eroded suture-line is shown as Figure 27.

*O. buarquianum* morphotypes are insensibly connected to *O. douglasi* morphotypes (Fig. 3A–B). Typically (Fig. 25A–B), this morphotype is strongly compressed, with 22–32 broad, low, weak ribs per whorl, and an indistinct umbilical wall. It is perhaps worth noting that it is a 'law' of polytypic ammonite species that the compressed variants are weakly ornamented, and that strength of ornament increases with inflation.

Two examples, SAM-PCA 3603–3604, show the rather dense, straight ribs of *O. salasi* Young, becoming projected near the venter (Fig. 19D–E), and are connected by intermediates to *O. douglasi*. These *O. salasi* variants show a low umbilical wall, broad convex flanks and a lanceolate whorl section, and were clearly rather evolute. Although the siphonal keel is not preserved, it certainly stood well above the venter. The dense, narrow, simple ribs are distinctly prorsiradiate and only very slightly flexuous, with at least twenty-two ribs per half whorl. The ribs swell slightly at the ventrolateral shoulders and curve strongly forward to meet the siphonal keel.

### Discussion

The Angolan material shows an insensible transition between *O. buarquianum* (White) and *O. douglasi* Knetchel and they are synonyms. It is unfortunate that White's (1887) species has priority because the *O. douglasi* morphotypes predominate. There is little to separate *O. packardi* Anderson from the *O. douglasi* morphotypes and it should also fall into synonymy. Several of the Angolan morphotypes are extremely close to *O. mirapelianum*, which is a broadly contemporaneous European species. However, until the population

structure of D'Orbigny's (1841) species is better understood, they are maintained as specifically distinct.

As already noted, there are *O. salasi* morphotypes within the Angolan material, which are connected by transitions to *O. douglasi* morphotypes, but they constitute only 4 per cent of the present material. Because of this, and because Young (1966) had over 300 specimens of *O. salasi* from the Key Valley Marl, the Texas form is maintained as a distinct species. Occasional *O. salasi* morphotypes within the Angolan population merely indicate that *O. salasi* is descended from an *O. buarquianum* ancestor by a major change in population structure. *Oxytropidoceras stenzeli* Young (1966: 93, pl. 22 (fig. 1), pl. 26 (fig. 1), pl. 34 (fig. 1), pl. 35 (fig. 4), figs 7h, 18c) is a Middle to low Upper Albian species that may not bear separation from *O. salasi*.

Collignon (1936) recorded *O. mirapelianum*, *O. colcanapi* Collignon, *O. boulei* Collignon, and *O. bravoense* Collignon (*non* Böse) from the same level and locality (Berambo) in the Middle Albian of Madagascar. His *O. colcanapi* is certainly very close to *O. buarquianum*, although the true affinities of the Madagascan material must await reassessment at the population level.

The species of *Oxytropidoceras* figured by Reymont (1955) from Nigeria are all poorly preserved juveniles with the simple, flexuous ribbing of *O. (Mirapelia)*. Additional and better preserved material is required for a satisfactory identification.

*Oxytropidoceras chihuahuaense* (Böse) (1910: 73, pl. 5 (figs 3–4), pl. 7 (figs 3–4), pl. 8 (figs 1–2)) was assigned by Young (1966) to *Venezolicer*, but the absence of a flank tubercle clearly places it in *O. (Mirapelia)*. The lectotype is the original of the specimen figured by Böse (1910, pl. 7 (figs 3–4), pl. 8 (fig. 1)), which shows rather rigid, slightly prorsiradiate single ribs that broaden ventrally and terminate in weak ventrolateral swellings. These features are within the range of variation of the Angolan material, but since Böse's species is, in Texas at least (Young 1966), dated as upper Middle Albian–lower Upper Albian, it is maintained as distinct.

*Oxytropidoceras involutum* Beurlen is based upon immature material that is very close to typical examples of this species. It is likely to prove a synonym of *O. buarquianum*.

### *Oxytropidoceras (Mirapelia) sergipense* (White, 1887)

Figs 11B, 18A–B, 23E, H, 28

*Ammonites sergipensis* White, 1887: 221, pl. 24 (figs 1–2).

*Pervinqueria sergipensis* (White) Maury, 1930: 294; 1936: 236, pl. 24 (figs 3–4).

?*Oxytropidoceras buarquianum* (White) Maury, 1936: 250, pl. 9 (fig. 1 only).

*Oxytropidoceras mauryae* Beurlen, 1952: 162, fig. 3.

*Dipoloceras sergipense* (White) Young, 1966: 20, 22, 24.

*Oxytropidoceras (Venezolicer) sergipense* (White) Renz, 1968: 630.

*Oxytropidoceras (Androiavites) mauryae* Beurlen, 1970: 468, pl. 6 (fig. 1).

*Non Mortoniceras sergipensis* (White) Beurlen, 1970: 468, pl. 6 (figs 2–4), fig. 19.

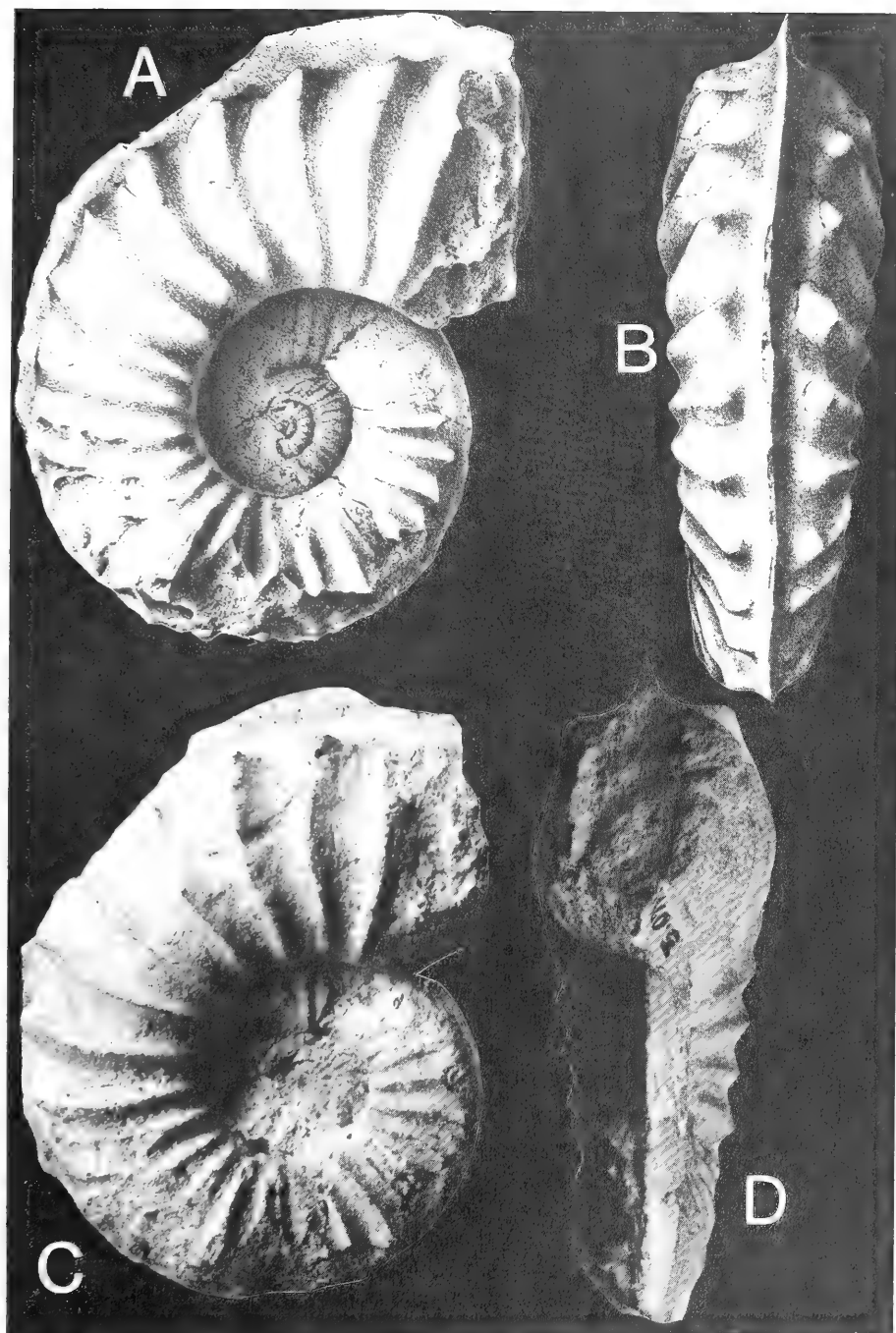


Fig. 28. *Oxytropidoceras (Mirapelia) sergipense* (White). A–B. White's (1887) original illustration. C–D. The original of White's illustration, herein selected as lectotype, in the Natural History Museum, Rio de Janeiro. Photo: P. Bengtson. Both  $\times 1$ .

### Material

Six specimens in the South African Museum, SAM-PCA3755, 3765, and 5444, together with three unnumbered examples. All are preserved as limestone steinkerns.

### Description

The shell is rather inflated and moderately evolute, with the outer whorl covering about 25 per cent of the preceding whorl. The umbilicus is moderately wide (32–35 % of the diameter) and fairly shallow, with steep umbilical walls and evenly rounded shoulders. Intercostally the whorl section is oval, compressed, whereas costally it is subquadrate (Fig. 23E, H). Ribs begin at the umbilical seam and, across the flank, are slightly flexuous and prorsiradiate, terminating in prominent ventrolateral tubercles that are projected forward on the venter. There are thirty ribs on the outer whorl. The flanks are convex and converge slightly towards the venter, with maximum width about one-third of the way up the flanks. The keel is not preserved but must have stood high above the venter.

### Discussion

The confusion surrounding the generic status of White's (1887) species is evident from the synonymy. It is, indeed, very close to *Venezoliceras*, as noted by Renz (1968), but lacks the development of flank tubercles. On the other hand, it is also very similar to the more inflated *Oxytropidoceras douglasi* morphotypes with which it coexists, differing only in the very pronounced development of ventrolateral tubercles. In gross morphology and stratigraphic position, therefore, *O. (M.) sergipense* is intermediate between *O. (Mirapelia)* and *Venezoliceras*. Phylogenetically, it is perhaps worth noting that Young (1966) assigned this species to *Dipoloceras*, while Spath (1932) referred such typical *O. (Mirapelia)* species as *O. evansi* (Spath), and even *O. mirapelianum* (d'Orbigny) itself, to *Dipoloceras*. As such, it seems likely that the origins of *Dipoloceras* also lie in *O. (Mirapelia)* of the *sergipense* group.

*Oxytropidoceras (Androiavites) mauryae* Beurlen shows the same prominent ventrolateral tubercles and simple ribbing as *O. (M.) sergipense* and is here regarded as conspecific. The specimen figured by Beurlen (1970) is an Upper Albian *Mortoniceras (Angolaites)* with distinct umbilical and doubled ventrolateral tubercles.

*Oxytropidoceras (Mirapelia) sergipense* (White) is very close to certain species of *Mojsisoviczia* (Figs 20, 22B–C), from which it is readily distinguished by its greater compression and the absence of a lateral tubercle. In this respect the figured but undescribed material of *Mojsisoviczia* aff. *delaruei* (d'Orbigny) (Spath 1930: 61, pl. 9 (figs 13, 16)) closely resembles the present species, since a lateral tubercle is not evident in the illustrations. It seems reasonable to assume that the ancestry of *Mojsisoviczia* lies in *O. (Mirapelia)* of the *sergipense* group.

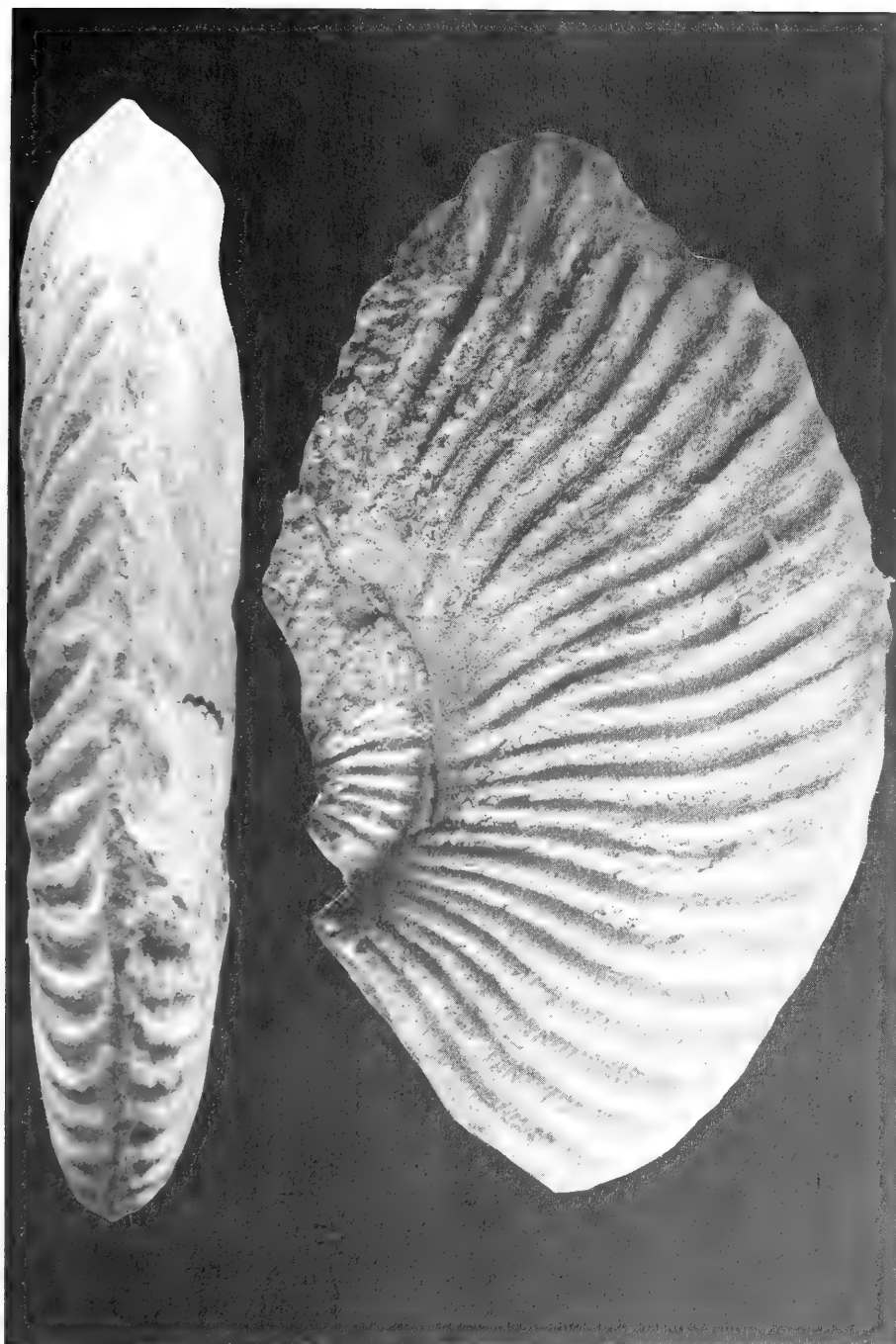


Fig. 29. *Oxytropidoceras (Oxytropidoceras) boesei* Knechtel, SAM-PCA5452,  $\times 1$ .

*Oxytropidoceras (Oxytropidoceras) boesei* Knechtel, 1947

Figs 29–31A

*Oxytropidoceras (Manuaniceras) bösei* Knechtel, 1947: 109, pl. 27 (fig. 1).? *Manuaniceras uddeni* Young, 1966: 95, pl. 12 (figs 1, 4–5), pl. 13 (fig. 1), pl. 16 (figs 1, 3), figs 10b, 12b.*Material*

Three specimens, SAM-PCA5452–5453 and an unnumbered example in the South African Museum, preserved as limestone steinkerns.

*Description*

The Angolan material comprises several rather eroded internal moulds of body chamber fragments that, because of scaphitoid-uncoiling of the body chamber, appear to be more evolute (umbilicus 33% of the diameter) than the earlier growth stages actually were. The shell is strongly compressed, with a lanceolate whorl section (Fig. 30) and broad, convex flanks. Maximum width is about one-third of the way up the flanks. Although the siphonal keel is not preserved, it certainly stood well above the venter.

Ornament comprises dense, flexuous flank ribs that arise at, or close to, the umbilical seam and pass forward across the flanks. The ribs are narrower than the interspaces with a steep, concave, adoral surface and a sloping, convex, adapical face. They frequently bifurcate close to the umbilical seam, but only rarely at, or above, mid-flank. There are 8–9 ribs along the venter in a distance equal to the whorl height. There is no sign of bullae on any part of the flanks.

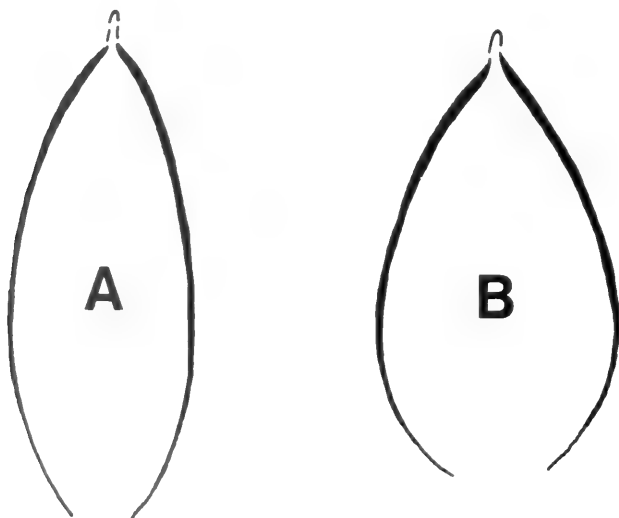


Fig. 30. *Oxytropidoceras (Oxytropidoceras) boesei* Knechtel. Whorl sections. A. SAM-PCA5452. B. SAM-PCA5453.  $\times 1$ .



### Discussion

*Oxytropidoceras (Manuaniceras) boesei* Knetchel was based upon fragmentary material, without proper description, which shows no features to distinguish it from the Angolan material. The Texas species, *Manuaniceras uddeni* Young, which coexists with *O. (M.) buarquianum*, is very close to the present material. It is said to differ in the presence of a few very weak bullae at the umbilical shoulder, and sometimes also at the point of bifurcation or intercalation of a rib. Since the presence of such bullae is likely to be markedly affected by state of preservation, *M. uddeni* is probably a synonym of *O. boesei*.

*Oxytropidoceras roissyanum* (d'Orbigny) (Fig. 5A–B) is difficult to compare with the present species because the known material of both species represents vastly different growth stages. So far as can be judged, *O. roissyanum* is much more inflated, with stronger, more robust ribs, about as wide as the interspaces, and does not show bifurcation above the umbilical shoulder. *Oxytropidoceras applanatus* Collignon (1936: 183, pl. 16, (fig. 5)) is doubtfully distinct from *O. roissyanum*.

*Oxytropidoceras manuanense* (Spath) (1921: 281, pl. 25 (fig. 1)) (Fig. 31B) resembles the present species in lacking flattened ribs, but differs in that nearly all ribs bifurcate at some level on the flanks and in the effacement of ribbing in maturity.

*Oxytropidoceras cantianum* Spath (Fig. 14F–G) is an Upper Albian (*Dipoloceras cristatum* Zone) species that is based upon a juvenile showing strongly branching ribs. It closely approaches the example figured by Young (1966, pl. 7 (figs 4, 6)) as 'probably *M. carbonarium* (Gabb)', and Owen (1971) seems to be right in regarding it a synonym of *O. carbonarium*.

*Oxytropidoceras aroeirium* Maury (1936: 251, pl. 26, (fig. 7)) is, to judge from the description, very close to the present species. Unfortunately, the writer has not seen the illustrations of this species and thus further comment is unjustified.

### AGE OF THE PRESENT FAUNA

*Douvilleiceras mammillatum* (Schlotheim) has long been considered diagnostic of the uppermost biostratigraphic zone in the Lower Albian, to which it gives its name. Thus, the rich *Douvilleiceras* faunas from the rest of the world are almost invariably assigned to the zone of *D. mammillatum* and considered to be of late early Albian age. Despite the fact that *Douvilleiceras* has long been known to persist into the Middle Albian (Casey 1962; Owen 1971; Destombes *et al.* 1977), including *D. mammillatum* itself (Fig. 12E–F), this has never been a problem in the boreal realm where the incoming of *Hoplites* forms an easily recognizable and diagnostic base to the Middle Albian. Outside the hoplitinid faunal province, however, no such aid is available and the problem of

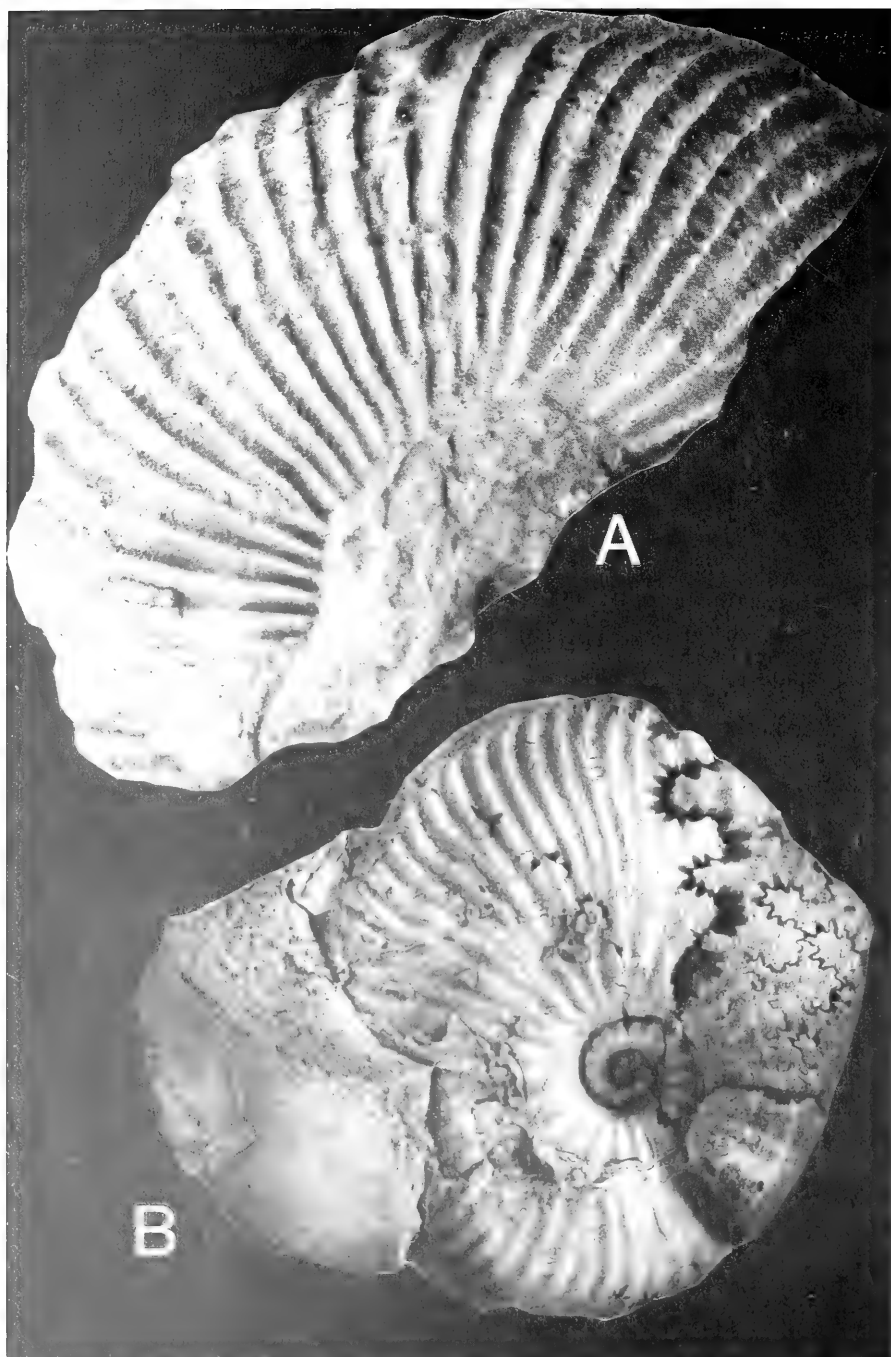


Fig. 31. A. *Oxytropidoceras* (*Oxytropidoceras*) *boesei* Knechtel, SAM-PCA5453. B. *Oxytropidoceras* (*Oxytropidoceras*) *man-uanense* (Spath), SAM-2726, a paratype.

the Lower–Middle Albian boundary is far more complex. These problems led Young (1966) to suggest that *Lyelliceras* may date a younger level in South America than it does in Europe. Contrary to Young's (1966) opinion, however, the writer feels that *Lyelliceras* is critical to the problem of the Lower–Middle Albian boundary, in view of its widespread geographic distribution and common association with hoplinitids in the boreal realm.

In the Anglo-Paris basin, the type region for the Albian Stage, Owen (1971) recognized the following subdivisions of the Middle Albian:

<i>Euhoplites lautus</i> Zone	{	<i>Anahoplites daviesi</i> Subzone
	{	<i>Euhoplites nitidus</i> Subzone
<i>Euhoplites loricatus</i> Zone	{	<i>Euhoplites meandrinus</i> Subzone
	{	<i>Mojsisoviczia subdelaruei</i> Subzone
	{	<i>Dimorphoplites niobe</i> Subzone
	{	<i>Anahoplites intermedius</i> Subzone
<i>Hoplites dentatus</i> Zone	{	<i>Hoplites spathi</i> Subzone
	{	<i>Lyelliceras lyelli</i> Subzone
	{	<i>Isohoplites eodentatus</i> Subzone

In the Anglo-Paris basin, *Douvilleiceras inaequinodum* (Quenstedt) does not range above the *eodentatus* Subzone, while *Oxytropidoceras* (*Mirapelia*) *mirapelianum* (d'Orbigny) is a constituent of the *Hoplites spathi* Subzone (Owen 1971). Moreover in Peru, Benavides-Cáceres (1956) records *Oxytropidoceras 'douglasi'* in association with *Lyelliceras lyelli* (d'Orbigny). There can also be little doubt that the Angolan *Oxytropidoceras* fauna is strictly contemporaneous with those from the Texan zone of *O. salasi*, and it is certainly also represented in Brazil (White 1887; Maury 1936; Beurlen 1970). As such, the *Lyelliceras lyelli* Subzone is represented in the south-central Atlantic region by *Oxytropidoceras buarquianum* (White) and/or *Lyelliceras lyelli* (d'Orbigny), and their associated species. The underlying beds in Angola, dominated by *Douvilleiceras mammillatum aequinodum* (Quenstedt) and *D. inaequinodum* (Quenstedt), are thus certainly to be correlated with the *Isohoplites eodentatus* Subzone of the Anglo-Paris basin. It is perhaps worthy of note that *O. mirapelianum* (d'Orbigny) is a replacement name for *Ammonites cristatus* Quenstedt (*non* DeLuc) and, as such, Quenstedt (1849) was dealing with a faunal horizon closely comparable, and probably strictly contemporaneous, with those at Dombé Grande.

As such, the Dombé Grande faunas are here regarded as of basal Middle Albian age, to be correlated with the *Isohoplites eodentatus* and *Lyelliceras lyelli* Subzones of the European succession. The absence of lyelliceratids and brancoceratids is enigmatic. Since *Oxytropidoceras* species are relatively long-lived, the *Oxytropidoceras* (*M.*) *buarquianum* faunule may prove to range somewhat higher. The fact that many of the *Douvilleiceras* assemblages from around the world could now prove to be of basal Middle Albian age suggests

that the late early Albian eustatic transgression (Cooper 1977) might actually have peaked during the earliest Middle Albian.

### SUMMARY

Two Middle Albian faunules rich in numbers, but poor in species, are described from the vicinity of Dombe Grande, Angola. The lower faunule, which is correlated with the *Isohoplites eodentatus* Subzone of the European sequence has yielded *Douvilleiceras mammillatum aequinodum* (Quenstedt), *D. inaequinodum* (Quenstedt), *D. variable* Tavani and *Puzosia bistricta* (White). The upper faunule has yielded *Oxytropidoceras* (*Oxytropidoceras*) *boesei* Knechtel, *O. (M.) buarquianum* (White), *O. (M.) sergipense* (White), *Puzosia bistricta* (White), *Protanisoceras* sp. and a *Hamites* sp., and is to be correlated with the *Lyelliceras lyelli* Subzone of the Anglo-Paris basin. Mojsisoviciinid phylogeny is discussed, and the following new taxa introduced: *Mortoniceratoides* gen. nov., *Oxytropidoceras* (*Mirapelia*) subgen. nov. and *O. (Benavidesites)* subgen. nov., while *Manuaniceras* is included in the synonymy of *Oxytropidoceras* s.s.

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6. SYSTEMATIC papers must conform to the *International code of zoological nomenclature* (particularly Articles 22 and 51).

Names of new taxa, combinations, synonyms, etc., when used for the first time, must be followed by the appropriate Latin (not English) abbreviation, e.g. gen. nov., sp. nov., comb. nov., syn. nov., etc.

An author's name when cited must follow the name of the taxon without intervening punctuation and not be abbreviated; if the year is added, a comma must separate author's name and year. The author's name (and date, if cited) must be placed in parentheses if a species or subspecies is transferred from its original genus. The name of a subsequent user of a scientific name must be separated from the scientific name by a colon.

Synonymy arrangement should be according to chronology of names, i.e. all published scientific names by which the species previously has been designated are listed in chronological order, with all references to that name following in chronological order, e.g.:

#### Family Nuculanidae

*Nuculana (Lembulus) bicuspidata* (Gould, 1845)

Figs 14–15A

*Nucula (Leda) bicuspidata* Gould, 1845: 37.

*Leda plicifera* A. Adams, 1856: 50.

*Laeda bicuspidata* Hanley, 1859: 118, pl. 228 (fig. 73). Sowerby, 1871: pl. 2 (fig. 8a–b).

*Nucula largillierii* Philippi, 1861: 87.

*Leda bicuspidata*: Nickles, 1950: 163, fig. 301; 1955: 110. Barnard, 1964: 234, figs 8–9.

Note punctuation in the above example:

comma separates author's name and year

semicolon separates more than one reference by the same author

full stop separates references by different authors

figures of plates are enclosed in parentheses to distinguish them from text-figures

dash, not comma, separates consecutive numbers

Synonymy arrangement according to chronology of bibliographic references, whereby the year is placed in front of each entry, and the synonym repeated in full for each entry, is not acceptable.

In describing new species, one specimen must be designated as the holotype; other specimens mentioned in the original description are to be designated paratypes; additional material not regarded as paratypes should be listed separately. The complete data (registration number, depository, description of specimen, locality, collector, date) of the holotype and paratypes must be recorded, e.g.:

#### Holotype

SAM-A13535 in the South African Museum, Cape Town. Adult female from mid-tide region, King's Beach Port Elizabeth (33° 51' S 25° 39' E), collected by A. Smith, 15 January 1973.

Note standard form of writing South African Museum registration numbers and date.

## 7. SPECIAL HOUSE RULES

### Capital initial letters

- The Figures, Maps and Tables of the paper when referred to in the text  
e.g. '... the Figure depicting *C. namacolus* ...'; '... in *C. namacolus* (Fig. 10) ...'
- The prefixes of prefixed surnames in all languages, when used in the text, if not preceded by initials or full names  
e.g. Du Toit but A. L. du Toit; Von Huene but F. von Huene
- Scientific names, but not their vernacular derivatives  
e.g. Therocephalia, but therocephalian

Punctuation should be loose, omitting all not strictly necessary

Reference to the author should be expressed in the third person

Roman numerals should be converted to arabic, except when forming part of the title of a book or article, such as

'Revision of the Crustacea. Part VIII. The Amphipoda.'

Specific name must not stand alone, but be preceded by the generic name or its abbreviation to initial capital letter, provided the same generic name is used consecutively.

Name of new genus or species is not to be included in the title: it should be included in the abstract, counter to Recommendation 23 of the Code, to meet the requirements of Biological Abstracts.



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LOWER CRETACEOUS  
(MIDDLE ALBIAN)  
AMMONITES FROM  
DOMBE GRANDE, ANGOLA